



**Global, regional, and national incidence, prevalence, and years lived with disability for
310 diseases and injuries, 1990-2015
a systematic analysis for the Global Burden of Disease Study 2015**

Global, regional, and national incidence, prevalence, and years lived with disability for 310 diseases and injuries, 1990–2015: a systematic analysis for the Global Burden of Disease Study 2015

GBD 2015 Disease and Injury Incidence and Prevalence Collaborators*



Background Non-fatal outcomes of disease and injury increasingly detract from the ability of the world's population to live in full health, a trend largely attributable to an epidemiological transition in many countries from causes affecting children, to non-communicable diseases (NCDs) more common in adults. For the Global Burden of Diseases, Injuries, and Risk Factors Study 2015 (GBD 2015), we estimated the incidence, prevalence, and years lived with disability for diseases and injuries at the global, regional, and national scale over the period of 1990 to 2015.

Methods We estimated incidence and prevalence by age, sex, cause, year, and geography with a wide range of updated and standardised analytical procedures. Improvements from GBD 2013 included the addition of new data sources, updates to literature reviews for 85 causes, and the identification and inclusion of additional studies published up to November, 2015, to expand the database used for estimation of non-fatal outcomes to 60 900 unique data sources. Prevalence and incidence by cause and sequelae were determined with DisMod-MR 2.1, an improved version of the DisMod-MR Bayesian meta-regression tool first developed for GBD 2010 and GBD 2013. For some causes, we used alternative modelling strategies where the complexity of the disease was not suited to DisMod-MR 2.1 or where incidence and prevalence needed to be determined from other data. For GBD 2015 we created a summary indicator that combines measures of income per capita, educational attainment, and fertility (the Socio-demographic Index [SDI]) and used it to compare observed patterns of health loss to the expected pattern for countries or locations with similar SDI scores.

Findings We generated 9·3 billion estimates from the various combinations of prevalence, incidence, and YLDs for causes, sequelae, and impairments by age, sex, geography, and year. In 2015, two causes had acute incidences in excess of 1 billion: upper respiratory infections (17·2 billion, 95% uncertainty interval [UI] 15·4–19·2 billion) and diarrhoeal diseases (2·39 billion, 2·30–2·50 billion). Eight causes of chronic disease and injury each affected more than 10% of the world's population in 2015: permanent caries, tension-type headache, iron-deficiency anaemia, age-related and other hearing loss, migraine, genital herpes, refraction and accommodation disorders, and ascariasis. The impairment that affected the greatest number of people in 2015 was anaemia, with 2·36 billion (2·35–2·37 billion) individuals affected. The second and third leading impairments by number of individuals affected were hearing loss and vision loss, respectively. Between 2005 and 2015, there was little change in the leading causes of years lived with disability (YLDs) on a global basis. NCDs accounted for 18 of the leading 20 causes of age-standardised YLDs on a global scale. Where rates were decreasing, the rate of decrease for YLDs was slower than that of years of life lost (YLLs) for nearly every cause included in our analysis. For low SDI geographies, Group 1 causes typically accounted for 20–30% of total disability, largely attributable to nutritional deficiencies, malaria, neglected tropical diseases, HIV/AIDS, and tuberculosis. Lower back and neck pain was the leading global cause of disability in 2015 in most countries. The leading cause was sense organ disorders in 22 countries in Asia and Africa and one in central Latin America; diabetes in four countries in Oceania; HIV/AIDS in three southern sub-Saharan African countries; collective violence and legal intervention in two north African and Middle Eastern countries; iron-deficiency anaemia in Somalia and Venezuela; depression in Uganda; onchocerciasis in Liberia; and other neglected tropical diseases in the Democratic Republic of the Congo.

Interpretation Ageing of the world's population is increasing the number of people living with sequelae of diseases and injuries. Shifts in the epidemiological profile driven by socioeconomic change also contribute to the continued increase in years lived with disability (YLDs) as well as the rate of increase in YLDs. Despite limitations imposed by gaps in data availability and the variable quality of the data available, the standardised and comprehensive approach of the GBD study provides opportunities to examine broad trends, compare those trends between countries or subnational geographies, benchmark against locations at similar stages of development, and gauge the strength or weakness of the estimates available.

Funding Bill & Melinda Gates Foundation.

Copyright © The Author(s). Published by Elsevier Ltd. This is an Open Access article under the CC BY license.

Lancet 2016; 388: 1545–602

This online publication has been corrected. The corrected version first appeared at thelancet.com on January 5, 2017

See Editorial page 1447

See Comment pages 1448 and 1450

*Collaborators listed at the end of the Article

Correspondence to:
Prof Theo Vos, Institute for Health Metrics and Evaluation, Seattle, WA 98121, USA
tvos@uw.edu

Introduction

Although substantial progress has been made toward reducing mortality and extending life expectancy throughout the world over the past few decades, the epidemiological transition is manifest in the growing importance of non-fatal diseases, outcomes, and injuries which pose, partly as a consequence of decreasing death rates, a rising challenge to the ability of the world's population to live in full health. Complementing information on deaths by age, sex, cause, geography, and time with equally detailed information on disease incidence, prevalence, and severity is key to a balanced debate in health policy. For this reason, the Global Burden of Disease (GBD) Study uses the disability-adjusted life-year (DALY), combining years of life lost (YLLs) due to mortality and years lived with disability (YLDs) in a single metric. One DALY can be thought of as one lost year of healthy life. The sum of DALYs in a population can be thought of as the gap between the population's present health status and an ideal situation where the entire population lives to an advanced age, free of disease. Assessments of how different diseases lead to multimorbidity and reductions in functional health status are important for both health system planning¹ and a broader range of social policy issues such as the appropriate age for retirement in some countries.^{2,3} Many challenges in making standardised estimates of non-fatal health outcomes are similar to those affecting mortality estimates (including variations in case definitions, data collection methods, variable quality of data collection, conflicting data, and missing data) but are compounded by more sparse and varied data sources, the need to characterise each disease by its disabling sequelae or consequence(s), and the need to quantify the severity of these consequences. The standardised approach of the annual GBD updates addresses these measurement problems to enhance comparability between causes by geography and over time.

The estimates from GBD 2013 drew attention to large increases in the number of YLDs over the previous decade, whereas rates of YLDs for most causes remained stable or showed only small decreases.⁴ The GBD 2013 assessment largely attributed increases in the number of YLDs to musculoskeletal disorders, mental and substance use disorders, neurological disorders, and chronic respiratory diseases, as well as population growth and ageing. GBD 2013 also brought attention to increased differences in trends between mortality and morbidity for many causes. YLDs as a proportion of DALYs increased globally, a manifestation of the continuing epidemiological transition in low-income and middle-income countries. Decreases in mortality from diseases such as pneumonia, diarrhoea, maternal and neonatal disorders, and an absence of progress in reducing YLD rates continued to drive a transition toward a greater global number of YLDs.

Along with broad recognition that data from some regions were sparse and that more and higher quality data in general would probably improve estimation, useful debates on the GBD results have been published. These debates have focused on the analysis or presentation of individual diseases, such as changes over time in GBD estimates of dementia,⁵ the accuracy of HIV incidence estimates,^{6,7} the absence of sepsis as a disease,^{8,9} the quality of some cancer registry data,¹⁰ and the absence of mental disorders as sequelae of neglected tropical diseases.¹¹ The GBD empirical approach to measuring the public's view of health state severity has generated substantial interest with questions about the relative importance of different dimensions of health,^{12,13} the quantification of health loss,^{14,15} and discussions of the transferability of judgments about relative health to conventional notions of disability and dependence.⁵ In each cycle of the GBD, we seek to improve the estimates, reflecting published and unpublished critique through the acquisition of new data, expansion of the network of collaborators, changes in how data are corrected for bias, advances in modelling techniques, and the targeted expansion of the GBD cause list.

The primary objective of this component of the GBD was to use all available data of sufficient quality to generate reliable and valid assessments of disease and injury sequelae incidence, prevalence, and YLDs for all 310 causes in the GBD cause hierarchy for 591 locations in the GBD study during 1990–2015. We describe the change over time and between populations in relation to where countries fall on the development continuum.¹⁶ Continuing efforts to improve data and code transparency are an important part of the GBD cycle. These results thus supersede any previous publications about the GBD on disease incidence, prevalence, and YLDs.

Methods

Overall approach

We estimated incidence and prevalence by age, sex, cause, year, and geography using a wide range of updated and standardised analytical procedures. The overall logic of our analytical approach is shown for the entire non-fatal estimation process in figure 1. The appendix provides a single source for detail of inputs, analytical processes, and outputs and methods specific to each cause. This study complies with the Guidelines for Accurate and Transparent Health Estimates Reporting (GATHER) recommendations (methods appendix pp 1, 608–10).¹⁷

Geographies in GBD 2015

The geographies included in GBD 2015 have been arranged into a set of hierarchical categories composed of seven super-regions and a further nested set of 21 regions containing 195 countries and territories. Eight additional subnational assessments were done for Brazil, China, India, Japan, Kenya, Saudi Arabia, South Africa, Sweden, and the USA (methods appendix pp 611–24). For this study we present data at the national and territory level.

See Online for appendix

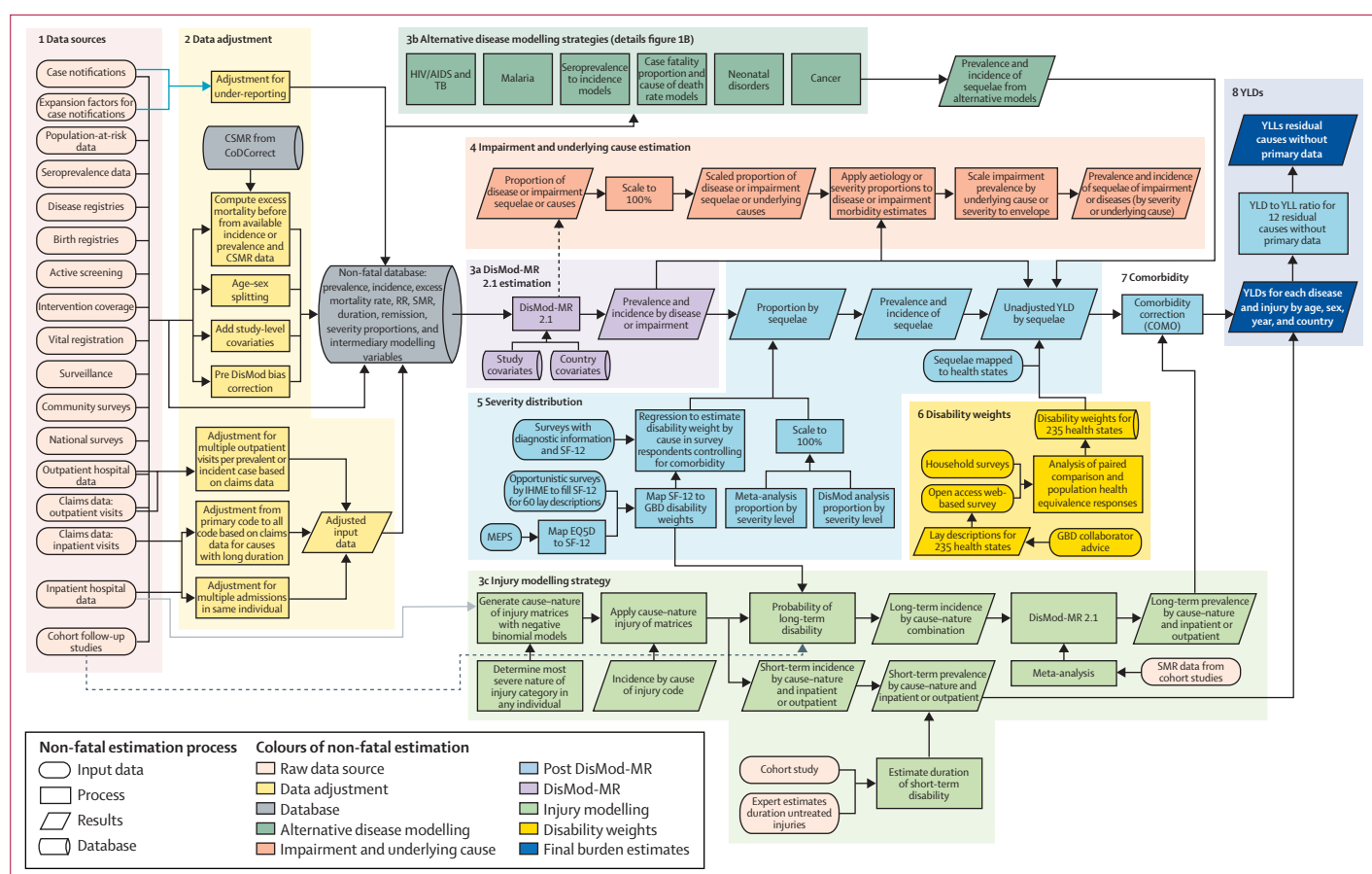


Figure 1: Analytical flow chart for the estimation of cause-specific YLDs by location, age, sex, and year for GBD 2015

Ovals represent data inputs, square boxes represent analytical steps, cylinders represent databases, and parallelograms represent intermediate and final results. The flow chart is colour-coded by major estimation component: raw data sources, in pink; data adjustments, in yellow; DisMod-MR 2.1 estimation, in purple; alternative modelling strategies, in light green; injury modelling strategy, in dark green; estimation of impairments and underlying causes, in brown; post-DisMod-MR and comorbidity correction, in blue; disability weights, in orange; and cause of death and demographic inputs, in grey. GBD=Global Burden of Disease. TB=tuberculosis. SF-12=Short Form 12 questions. MEPS=Medical Expenditure Panel Surveys. CSMR=cause-specific mortality rate. SMR=standardised mortality ratio. YLDs=years lived with disability. YLLs=years of life lost. IHME=Institute for Health Metrics and Evaluation.

List of causes and sequelae

The GBD cause and sequelae list is organised hierarchically (methods appendix 625–53). At Level 1 there are three cause groups: communicable, maternal, neonatal, and nutritional diseases (Group 1 diseases); non-communicable diseases; and injuries. These Level 1 aggregates are subdivided at Level 2 of the hierarchy into 21 cause groupings. The disaggregation into Levels 3 and 4 contains the finest level of detail for causes captured in GBD 2015. Sequelae of diseases and injuries are organised at Levels 5 and 6 of the hierarchy. The finest detail for all sequelae estimated in GBD is at Level 6 and is aggregated into summary sequelae categories (Level 5) for causes with large numbers of sequelae. Sequelae in GBD are mutually exclusive and collectively exhaustive, and thus our YLD estimates at each level of the hierarchy sum to the total of the level above. Prevalence aggregations are estimated at the level of individuals who might have more than one sequela or disease and therefore are not additive.

The cause and sequelae list was expanded based upon feedback after the release of GBD 2013 and input from GBD 2015 collaborators. Nine causes for which non-fatal outcomes are estimated were added: Ebola virus disease, motor-neuron disease, environmental heat and cold exposure, four subtypes of leukaemia, and two subtypes of non-melanoma skin cancer (methods appendix pp 625–53). The incorporation of these changes expanded the cause list from the 301 causes with non-fatal estimates examined in GBD 2013, to 310 causes with non-fatal estimates and from 2337 to 2619 unique sequelae at Level 6 of the hierarchy. At the newly created Level 5 of the hierarchy there were 154 summary sequela categories. The methods appendix (pp 654–61) provides a list of International Classification of Diseases version 9 (ICD-9) and version 10 (ICD-10) codes used in the extraction of hospital and claims data, mapped to GBD 2015 non-fatal causes, impairments, and nature of injury categories.

Period of analysis

A complete set of age-specific, sex-specific, cause-specific, and geography-specific incidence and prevalence numbers and rates were computed for the years 1990, 1995, 2000, 2005, 2010, and 2015. In this study we focus on trends for main and national results over the past decade, from 2005 to 2015, together with more detailed results for 2015. Online data visualisations at vizhub provide access to results for all GBD metrics.

Non-fatal modelling strategies vary substantially between causes. Figure 1 outlines the general process of non-fatal outcome estimation from data inputs to finalisation of YLD burden results; step 3b of that process identifies alternative modelling approaches used for specific causes (methods appendix pp 603, 604). The starting point for non-fatal estimation is the compilation of data sources identified through systematic analysis and extractions based on predetermined inclusion and exclusion criteria (methods appendix p 603). As part of the inclusion criteria, we defined disease-specific or injury-specific reference case definitions and study methods, as well as alternative allowable case definitions and study methods which were adjusted for if we detected a systematic bias. We used 15 types of primary data sources representing disease prevalence, incidence, mortality risk, duration, remission, or severity in the estimation process (oval shapes in figure 1).

Data sources

For this iteration of the study, we updated data searches through systematic data and literature reviews for 85 causes published up to Oct 31, 2015. For other causes, input from GBD collaborators resulted in the identification and inclusion of a small number of additional studies published after January, 2013. Data were systematically screened from household surveys archived in the Global Health Data Exchange, sources suggested to us by in-country experts, and surveys identified in major multinational survey data catalogues and Ministry of Health and Central Statistical Office websites. Case notifications reported to WHO were updated up to and including 2015. Citations for all data sources used for non-fatal estimation in GBD 2015 are provided in searchable form through a new web tool. A description of the search terms used for cause-specific systematic reviews, inclusion and exclusion criteria, and the preferred and alternative case definitions and study methods are detailed by cause in the methods appendix (pp 26–601).

Hospital inpatient data were extracted from 284 country-year and 976 subnational-year combinations from 27 countries in North America, Latin America, Europe, and New Zealand. Outpatient encounter data were available from the USA, Norway, Sweden, and Canada for 48 country-years. For GBD 2015, we also accessed aggregate data derived from claims information in a database of US private and public insurance schemes for the years 2000, 2010, and 2012. From the linked

claims data, we generated several correction factors to account for bias in health service encounter data from elsewhere, which were largely available to us aggregated by ICD code and by primary diagnosis only. First, for chronic disorders, we estimated the ratio between prevalence from primary diagnoses and prevalence from all diagnoses associated with a claim. Second, we used the claims data to generate the average number of outpatient visits per disorder. Similarly, we generated per person discharge rates from hospital inpatient data in the USA and New Zealand, the only sources with unique patient identifiers available for GBD 2015.

In GBD 2013, we calculated a geographical and temporal data representativeness index (DRI) of non-fatal data sources for each cause or impairment. The DRI represents the fraction of countries for which any incidence, prevalence, remission, or mortality risk data were available for a cause. This metric quantifies data availability, not data quality. The overall DRI and period-specific DRI measures for each cause and impairment are presented in the methods appendix (pp 662–68). DRI ranged from 90% for nine causes, including tuberculosis and measles, to less than 5% for acute hepatitis C and the category of other exposures to mechanical forces. Required case reporting resulted in high DRI values for notifiable infectious diseases; the network of population-based registries for cancers resulted in a DRI of above 50%. DRI values ranged from 6.1% in North Korea to 91.3% in the USA. Many high-income countries, as well as Brazil, India, and China, had DRI values above 63%; data availability was low in several countries, including Equatorial Guinea, Djibouti, and South Sudan.

Non-fatal disease models

In addition to the corrections applied to claims and hospital data, a number of other adjustments were applied including age–sex splitting, bias correction, adjustments for under-reporting of notification data, and computing expected values of excess mortality. In GBD 2013, we estimated expected values of excess mortality from prevalence or incidence and cause-specific mortality rate data for a few causes only, including tuberculosis and chronic obstructive pulmonary disease. In order to achieve greater consistency between our cause of death and non-fatal data, we adopted this strategy systematically for GBD 2015. We matched every prevalence data point (or incidence datapoint for short duration disorders) with the cause-specific mortality rate value corresponding to the age range, sex, year, and location of the datapoint. The ratio of cause-specific mortality rate to prevalence is conceptually equivalent to an excess mortality rate.

To estimate non-fatal health outcomes in previous iterations of GBD, most diseases and impairments were modelled in DisMod-MR, a Bayesian meta-regression tool originally developed for GBD 2010 (step 3a in figure 1).¹⁸ DisMod-MR was designed to address statistical challenges in estimation of non-fatal health outcomes, and for

For data visualisations at vizhub see <http://vizhub.healthdata.org/gbd-compare>

For Global Health Data Exchange see <http://ghdx.healthdata.org>

For data in GBD 2015 see <http://ghdx.healthdata.org/global-burden-disease-study-2015>

synthesis of often sparse and heterogeneous epidemiological data. For GBD 2015, the computational engine of DisMod-MR 2.1 remained unchanged, but we substantially rewrote the code that organises the flow of data and settings at each level of the analytical cascade. The sequence of estimation occurs at five levels: global, super-region, region, country, and where applicable, subnational locations (appendix pp 611–24). At each level of the cascade, the DisMod-MR 2.1 computational engine enforces consistency between all disease parameters. For GBD 2015, we generated fits for the years 1990, 1995, 2000, 2005, 2010, and 2015. We log-linearly interpolated estimates for the intervening years in each 5-year period. Greater detail on DisMod-MR 2.1 is available at Global Health Data Exchange and the methods appendix (pp 7–11).

In previous iterations of GBD, custom models were created for a short list of causes for which the compartment model underpinning DisMod (susceptible, diseased, and dead) was insufficient to capture the complexity of the disease or for which incidence and prevalence needed to be derived from other data. Step 3b of figure 1 describes the development of custom models with greater detail shown in the methods appendix figure 1B (p 604, and for associated write-ups pp 26–601) for HIV/AIDS, tuberculosis, malaria, cancer, neonatal disorders, infectious diseases for which we derived incidence from seroprevalence data, and infectious diseases for which we derived incidence from cause of death rates and pooled estimates of the case fatality proportion.

In GBD 2013, we estimated the country–age–sex–year prevalence of nine impairments (step 4 of figure 1). Impairments in GBD are disorders or specific domains of functional health loss that are spread across many GBD causes as sequelae and for which there are better data to estimate the occurrence of the overall impairment than for each sequela based on the underlying cause. Overall impairment prevalence was estimated with DisMod-MR 2.1 except for anaemia, for which spatiotemporal Gaussian Process regression methods were applied. We constrained cause-specific estimates of impairments, such as in the 19 causes of blindness, to sum to the total prevalence estimated for that impairment. Anaemia, epilepsy, hearing loss, heart failure, and intellectual disability were estimated at different levels of severity.

Severity distributions

In step 5, sequelae were further defined in terms of severity for 194 causes at Level 4 of the hierarchy (figure 1A). We generally followed the same approach for estimating the distribution of severity as in GBD 2013. For Ebola virus disease, we created a health state for the infectious disease episode with duration derived from average hospital admission times, and a health state for ongoing postinfection malaise and joint problems based on four follow-up studies^{19–22} from which we derived an average duration. The health states for the subtypes of leukaemia and non-melanoma skin cancer were the

same as the general cancer health states. For motor-neuron disease we accessed the Pooled Resource Open-Access ALS Clinical Trials (PROACT) database containing detailed information on symptoms and impairments for more than 8500 patients who took part in the trials.²³

Disability weights

We used the same disability weights as in GBD 2013 (see methods appendix pp 669–94 for a complete listing of the lay descriptions and values for the 235 health states used in GBD 2015).

Comorbidity

In step 7, we estimated the co-occurrence of different diseases by simulating 40 000 individuals in each geography–age–sex–year combination as exposed to the independent probability of having any of the sequelae included in GBD 2015 based on disease prevalence. We tested the contribution of dependent and independent comorbidity in the US Medical Expenditure Panel Surveys (MEPS) data, and found that independent comorbidity was the dominant factor even though there are well known examples of dependent comorbidity. Age was the main predictor of comorbidity such that age-specific microsimulations accommodated most of the required comorbidity correction. Taking dependent comorbidity into account changed the overall YLDs estimated in the MEPS data by only 2.5% (and ranging from 0.6% to 3.4% depending on age) in comparison to assuming independent comorbidity (methods appendix pp 18–20).²⁴

YLD computation

We report 95% uncertainty intervals (95% UI) for each quantity in this analysis using 1000 samples from the posterior distribution of prevalence and 1000 samples of the disability weight to generate 1000 samples of the YLD distribution. The 95% UI is reported as the 25th and 975th values of the distribution. We report significant changes in disease estimates between countries or over time if the change was noted in more than 950 of the 1000 samples computed for each result. For GBD 2015, we computed age-standardised prevalence YLD rates from the updated world population age standard developed for GBD 2013.²⁵ Less common diseases and their sequelae were included in 35 residual categories (methods appendix pp 695–97). For 22 of these residual categories, estimates were made from epidemiological data for incidence or prevalence. For 13 residual categories, we estimated YLDs by multiplying the residual YLL estimates by the ratio of YLDs to YLLs from the estimates for explicitly modelled Level 3 causes in the same disease category.

Socio-demographic Index

In GBD 2013, a sociodemographic status variable was computed based on a principal components analysis of income per capita, educational attainment, average age of the population, and the total fertility

For DisMod-MR 2.1 engine and the code see <http://ghdx.healthdata.org/global-burden-disease-study-2015>

	Incidence (thousands)		Percentage change (%)
	2005	2015	
Upper respiratory infections	15 624 257 (13 851 237–17 411 199)	17 230 659 (15 351 516–19 172 439)	10.3 (9.0 to 11.6)*
Diarrhoeal diseases	2 235 739 (2 139 050–2 348 407)	2 392 517 (2 301 101–2 503 094)	7.0 (6.0 to 8.0)*
Permanent caries	426 963 (371 216–484 332)	487 629 (423 507–552 895)	14.2 (13.1 to 15.4)*
Otitis media	429 820 (353 915–525 059)	471 027 (386 606–577 286)	9.6 (7.9 to 11.2)*
Lower respiratory infections	273 131 (257 286–288 078)	291 759 (276 244–307 004)	6.8 (5.6 to 8.1)*
Malaria	339 275 (261 417–447 605)	286 859 (219 712–377 332)	–15.4 (–23.0 to –8.1)*
Gastritis and duodenitis	185 250 (167 471–204 974)	213 729 (192 486–236 339)	15.4 (10.8 to 17.2%)*
Pyoderma	178 382 (172 199–184 147)	207 452 (200 498–213 936)	16.3 (15.6% to 17.0)*
Gonococcal infection	138 220 (107 106–184 385)	172 676 (129 731–235 737)	24.9 (17.9 to 31.0)*
Interstitial nephritis and urinary tract infections	127 380 (124 308–130 683)	152 295 (148 748–156 177)	19.6 (19.2 to 19.9)*
Varicella and herpes zoster	128 678 (124 298–133 090)	142 413 (137 804–147 181)	10.7 (10.0 to 11.4)*
Trichomoniasis	121 948 (104 825–141 284)	140 781 (121 207–163 163)	15.4 (14.5 to 16.5)*
Acute hepatitis A	109 609 (101 813–117 648)	114 212 (103 349–124 776)	4.2 (–7.2 to 17.0)
Hepatitis B	98 277 (86 507–112 527)	111 212 (97 410–126 251)	13.2 (–4.3 to 35.2)
Gallbladder and biliary diseases	88 215 (79 276–96 495)	104 322 (93 074–114 430)	18.3 (16.0 to 20.7)*
Peptic ulcer disease	83 388 (77 760–89 435)	87 410 (80 343–94 506)	4.8 (2.6 to 7.0)*
Dengue	32 749 (18 879–68 335)	79 609 (53 784–169 704)	143.1 (–0.3 to 564.7)
Other sense organ diseases	60 659 (58 721–62 579)	69 945 (67 856–72 080)	15.3 (14.6 to 16.0)*
Chlamydial infection	56 976 (45 489–70 839)	61 173 (48 871–76 698)	7.4 (5.5 to 9.4)*
Maternal abortion, miscarriage, and ectopic pregnancy	53 942 (43 630–67 168)	53 958 (43 224–67 417)	0.0 (–3.8 to 4.0)
Syphilis	43 515 (37 479–51 054)	45 413 (37 787–54 921)	4.4 (–0.3 to 8.1)
Deciduous caries	41 353 (28 723–58 131)	43 688 (29 630–62 543)	5.6 (2.1 to 8.1)*
Genital herpes	29 112 (25 131–33 432)	39 791 (35 569–44 572)	36.7 (32.4 to 41.6)*
Urolithiasis	17 875 (16 320–19 728)	22 080 (20 183–24 295)	23.5 (21.2 to 25.6)*
Cellulitis	17 312 (15 988–18 739)	21 211 (19 582–22 985)	22.5 (21.0 to 24.1)*
Maternal hypertensive disorders	20 416 (17 593–23 417)	20 731 (17 355–24 379)	1.5 (–3.0 to 6.3)
Acute hepatitis E	18 869 (17 340–20 580)	19 525 (18 011–21 273)	3.5 (2.2% to 4.7)*
Whooping cough	22 457 (17 322–28 268)	16 298 (12 599–20 445)	–27.4 (–29.5 to –25.2)*

(Table 1 continues on next page)

rate.²⁶ For GBD 2015, we excluded mean age of the population because it is directly affected by death rates. To improve interpretability for GBD 2015, we computed a Socio-demographic Index (SDI) similar to the computation of the human development index.²⁷ In the SDI, each component was weighted equally and rescaled from zero (for the lowest value observed during 1980–2015) to one (for the highest value observed) for income per capita and average years of schooling, and the reverse for the total fertility rate. The final SDI score was computed as the geometric mean of each of the components. SDI ranged from 0.060 in Mozambique in 1987 to 0.978 in District of Columbia, USA, in 2015.

Role of the funding source

The funder of the study had no role in study design, data collection, data analysis, data interpretation, or writing of the report. The corresponding author had full access to all the data in the study and had final responsibility for the decision to submit for publication.

Results

Global incidence and prevalence

We generated over 9.3 billion outcomes of incidence, prevalence, and YLDs for 310 diseases, injuries, and aggregate categories; 2619 unique and aggregate sequelae; nine impairments; 63 age–sex groups; 591 geographies; and 26 individual years from 1990 to 2015. Each of these 9.3 billion estimates was calculated 1000 times to determine uncertainty intervals. Here, we present key summary findings on global incidence of short duration diseases, global prevalence of long-term disorders, global prevalence of impairments, global numbers and rates of YLDs and changes from 2005 to 2015, global YLL and YLD rates of change, patterns of comorbidity, the expected changes in the composition of YLDs with SDI, and country findings of leading causes of YLDs.

Disorders of less than 3 months duration and injuries with incidence of more than 1 million cases per year in 2015 are listed in table 1. There were two disorders with incidence greater than 1 billion per year: upper respiratory infections (17.2 billion [95% UI 15.4–19.2 billion]), and diarrhoeal diseases (2.39 billion [2.30–2.50 billion]). A further 13 diseases and injuries caused between 100 million and 1 billion incident cases a year and 16 diseases and injuries had incident cases of between 10 million and 100 million per year (table 1).

The disease and injury sequelae with a duration of more than 3 months and a global prevalence of more than 1% in 2015 are presented in table 2, aggregated to the cause level. Prevalence for impairments is presented at the bottom of table 3. Eight out of 56 high-prevalence causes affected more than 10% of the world's population in 2015. A further 48 causes affected between 1% and 10% of the world's population (table 2). Although many of

these causes are not among the dominant causes of YLDs because of comparatively low average disability weights, some causes, such as headaches, gynaecological diseases, oral disorders, and skin diseases, put great demands on health system resources by their sheer numbers.

Anaemia was the most common of our nine impairments, affecting 2·36 billion (2·35–2·37 billion) people in 2015. The next most common impairments were hearing loss of greater than 20 dB (1·33 billion [1·26–1·40 billion]), vision loss (940 million [905–974 million]), developmental intellectual disability (153 million [114–191 million]), infertility (113 million [93·4–136 million]), heart failure (40·0 million [38·6–41·4 million]), and epilepsy (39·2 million [34·3–43·7 million]; table 3; see results appendix (pp 740–48) for the prevalence estimates of the underlying causes of these impairments). Iron deficiency was the cause of anaemia in more than half of all cases. Over 90% of hearing loss was classified as age-related or other hearing loss. The largest number of people with vision loss had uncorrected refraction error. Idiopathic developmental intellectual disability, idiopathic female infertility, and idiopathic epilepsy were the most common causes of their impairments. Ischaemic heart disease was the most common cause of heart failure.

Global causes of disability

Global trends in YLDs 2005 to 2015

GBD 2015 included the assessment of 2619 sequelae at Level 6 of the GBD cause hierarchy, including 1316 sequelae from injuries that contributed to the global burden of disability. Causes at Level 4 of the hierarchy that resulted in 30 million or more YLDs in 2015 included lower back pain, major depressive disorder, age-related and other hearing loss, and neck pain. Figure 2 compares the leading causes of global YLDs in 2005 and 2015, using the cause breakdowns at Level 3 of the GBD cause hierarchy. Among the ten leading causes of YLDs, iron-deficiency anaemia and depressive disorders switched ranks to positions three and four respectively, diabetes rose from the eighth to the sixth position, migraine dropped from position six to seven, and other musculoskeletal disorders dropped from rank seven to eight (figure 2).

Estimates of prevalence and YLDs at the global level for 2005 and 2015 for each cause are presented in table 3 (see results appendix pp 717–40 for full detail at the sequelae level). Prevalence and age-standardised YLDs for 21 Group 1 diseases decreased significantly and by more than 10%, including measles, African trypanosomiasis, diphtheria, lymphatic filariasis, and rabies.

Age-standardised YLD rates for all maternal causes and sequelae combined decreased between 2005 and 2015, whereas overall age-standardised YLDs for neonatal disorders increased by 3·11% (–0·32–6·79%) since 2005 to 144 YLDs per 100 000 (109–185 YLDs per 100 000) in 2015, and age-standardised YLD rates decreased and absolute numbers of cases increased for nutritional deficiencies.

	Incidence (thousands)		Percentage change (%)
	2005	2015	
(Continued from previous page)			
Typhoid fever	15 341 (13 454–17 417)	12 538 (10 887–14 283)	–18·3 (–20·8 to –15·6)*
Maternal sepsis and other maternal infections	11 938 (9457–15 125)	11 817 (9300–15 009)	–1·0 (–4·6 to 2·8)
Appendicitis	10 159 (9549 to 10 840)	11 619 (10 918 to 12 407)	14·4 (13·5 to 15·2)*
Pancreatitis	7160 (6 660–7 694)	8 902 (8 212–9 643)	24·3 (21·9 to 26·6)*
Maternal haemorrhage	8438 (6614–10 620)	8740 (6774–11 061)	3·6 (–0·3 to 7·6)
Other sexually transmitted diseases	7569 (6511–8741)	7656 (6583–8853)	1·2 (–0·1 to 2·4)
Ischaemic heart disease	6308 (5918–6716)	7287 (6798–7808)	15·5 (14·3 to 16·8)*
Maternal obstructed labour and uterine rupture	6830 (5202–8933)	6521 (4977–8588)	–4·5 (–9·1 to –0·3)*
Hepatitis C	4609 (4316 to 4918)	5394 (5032 to 5778)	17·0 (15·7 to 18·4)*
Ischaemic stroke	4651 (4397 to 4912)	5385 (5017 to 5726)	15·8 (13·4 to 18·4)*
Measles	15 610 (6699–31 357)	4652 (2072–9206)	–70·2 (–73·6 to –66·0)*
Paratyphoid fever	5580 (4498–6896)	4587 (3738–5601)	–17·8 (–22·6 to –12·7)*
Haemorrhagic stroke	3144 (2973–3307)	3583 (3336–3822)	14·0 (11·4 to 16·2)*
Paralytic ileus and intestinal obstruction	2664 (2494–2855)	3167 (2946–3406)	18·9 (16·9 to 20·7)*
Encephalitis	1489 (1380–1608)	1603 (1472–1750)	7·7 (5·9 to 9·5)*
Acute glomerulonephritis	1528 (1395–1672)	1534 (1394–1685)	0·4 (–2·6 to 3·1)

Data in parentheses are 95% UIs. *Percentage changes that are statistically significant.

Table 1: Global incidence of short duration (less than 3 months) sequelae in 2005 and 2015 for all ages and both sexes combined, with percentage change between 2005 and 2015 for level 4 causes with incidence greater than 1 million cases per year

Data in parentheses are 95% UIs. *Percentage changes that are statistically significant.

Table 1: Global incidence of short duration (less than 3 months) sequelae in 2005 and 2015 for all ages and both sexes combined, with percentage change between 2005 and 2015 for level 4 causes with incidence greater than 1 million cases per year

Age-standardised YLD rates attributable to hepatitis A, B, and C increased, and decreased for hepatitis E. The number of individuals with chronic hepatitis C infection increased from 121 million (108–133 million) in 2005 to 142 million (127–157 million) in 2015.

Changes in age-standardised YLDs and YLLs over time

In 2005, non-communicable diseases (NCDs) accounted for 23 of the leading 25 causes of age-standardised YLDs worldwide and 23 of the 25 leading causes in 2015 (figure 2). Although diabetes rose only two ranks in the list of leading cause of YLDs, from position eight to six between 2005 and 2015, the increase in age-standardised rate was 5·4% (3·2–7·5%). Musculoskeletal disorders occupied three of the leading 25 causes of disability in both 2005 and 2015; lower back and neck pain were the single largest cause with little change in their rates.

	Prevalence (thousands)		Percentage change (%)
	2005	2015	
Permanent caries	2 045 859 (1 909 845–2 170 355)	2 344 628 (2 193 751–2 488 741)	14.6 (13.7 to 15.5)*
Tension-type headache	1 306 390 (1 160 423–1 463 889)	1 505 892 (1 337 310–1 681 575)	15.3 (14.0 to 16.6)*
Iron-deficiency anaemia	1 456 387 (1 449 846–1 462 782)	1 477 531 (1 470 902–1 485 322)	1.5 (0.9 to 2.1)*
Age-related and other hearing loss	943 504 (886 325–995 403)	1 210 055 (1 140 224–1 274 665)	28.3 (27.4 to 29.2)*
Migraine	831 726 (755 991–918 968)	958 789 (872 109–1 055 631)	15.3 (14.0 to 16.6)*
Genital herpes	716 115 (626 987–817 387)	845 826 (736 724–968 066)	18.1 (16.4 to 19.9)*
Refraction and accommodation disorders	672 165 (649 969–694 024)	819 307 (789 917–848 059)	21.9 (20.5 to 23.2)*
Ascariasis	854 489 (790 000–924 895)	761 894 (682 558–861 031)	-10.8 (-22.5 to 2.5)
G6PD trait	663 704 (625 032–703 799)	728 549 (676 735–781 534)	9.8 (7.9 to 11.4)*
Acne vulgaris	605 008 (568 577–642 061)	632 741 (595 242–671 249)	4.6 (3.5 to 5.6)*
Other skin and subcutaneous diseases	492 883 (480 852–505 426)	605 036 (589 500–619 676)	22.8 (22.1 to 23.4)*
Deciduous caries	534 122 (449 525–635 866)	558 028 (462 649–669 027)	4.5 (2.3 to 6.1)*
Low back pain	460 164 (444 680–477 119)	539 907 (521 449–559 556)	17.3 (16.5 to 18.2)*
Periodontal diseases	428 784 (372 953–498 682)	537 506 (465 114–625 889)	25.4 (24.1 to 26.5)*
Fungal skin diseases	434 604 (395 512–475 904)	492 373 (448 951–538 232)	13.3 (12.5 to 14.1)*
Trichuriasis	473 399 (443 689–505 928)	463 652 (426 621–502 939)	-2.1 (-12.0 to 9.0)
Diabetes	333 325 (310 773–355 510)	435 328 (404 736–468 562)	30.6 (28.0 to 33.0)*
Premenstrual syndrome	391 207 (375 009–407 896)	430 697 (410 841–450 494)	10.1 (7.9 to 11.8)*
Hookworm disease	462 111 (430 885–495 596)	428 246 (394 486–468 292)	-7.3 (-16.9 to 3.5)
Sickle-cell trait	338 756 (318 736–378 582)	404 566 (381 223–448 155)	19.4 (18.3 to 20.3)*
Asthma	327 097 (296 406–358 060)	358 198 (323 134–393 466)	9.5 (7.6 to 11.6)*
Neck pain	295 532 (258 878–338 138)	358 007 (313 408–409 411)	21.1 (19.0 to 23.3)*
Hepatitis B	293 745 (284 478–303 036)	343 251 (330 541–357 195)	16.9 (15.3 to 18.4)*
Other musculoskeletal disorders	283 317 (254 135–315 519)	342 068 (305 431–385 147)	20.7 (17.5 to 24.0)*
Urolithiasis	259 567 (238 100–281 892)	318 763 (290 695–349 154)	22.8 (20.8 to 24.9)*
Thalassaemias trait	252 798 (247 008–259 972)	279 451 (272 819–287 357)	10.5 (10.1 to 11.0)*
Malaria	207 773 (186 530–230 942)	278 961 (240 158–320 921)	34.3 (26.8 to 42.3)*
Edentulism and severe tooth loss	216 473 (207 563–226 331)	275 619 (264 201–288 252)	27.3 (26.9 to 27.7)*
Anxiety disorders	232 597 (204 165–264 445)	267 202 (234 064–306 318)	14.9 (13.0 to 16.8)*

(Table 2 continues on next page)

However, age-standardised rates of YLDs increased for osteoarthritis 3.90% (3.00–4.83%) by 2015. Depressive disorders were the fourth leading cause of disability in 2005 and the third leading cause of disability in 2015, and age-standardised YLD rates associated with the disorder increased marginally (1.0% [0.5–1.5%]). Age-standardised rates of YLDs from alcohol use disorders decreased after 2005 (4.5% [2.3–6.4%]) whereas disability due to drug use disorders increased by 8.2% (6.2–10.2%).

In contrast with global trends for NCDs, both relative ranks and age-standardised YLD rates decreased for most injuries. Falls, which were the 13th leading cause of disability in 2005, dropped to the 15th rank, and age-standardised YLD rates decreased (8.58% [5.23–12.2%]). Other unintentional injuries decreased in global rank from 27th in 2005 to 34th in 2015, and age-standardised YLD rates decreased by 16.7% (15.7–17.9%).

The leading causes of disability varied considerably with age (figure 3). The leading cause in children younger than 5 years was iron-deficiency anaemia followed by skin diseases, protein-energy malnutrition, and diarrhoea. In older children, iron-deficiency anaemia, skin diseases, asthma, and mental health disorders such as conduct, autistic spectrum, and anxiety disorders were top ten causes of disability. In adolescents and young adults (aged between 15 and 39 years), iron-deficiency anaemia, skin diseases, depression, lower back and neck pain, and migraine led the rankings. Other mental health disorders such as anxiety disorders and schizophrenia were in the top ten causes in this age group. In middle-aged adults, musculoskeletal disorders dominated the top rankings followed by mental health disorders, especially depression. Diabetes and sense organ disorders were more prominent causes of disability in middle-age. In older adults (older than 65 years), sense organ disorders were the top-ranked cause of disability. Musculoskeletal disorders remained a dominant source of disability, and chronic obstructive pulmonary disease entered the top ten. In the oldest age groups, ischaemic heart disease and Alzheimer's and other dementias made their first appearance in the top ten.

We examined the trends in YLDs and YLLs in a scatterplot (figure 4). YLLs decreased for the majority of causes. For Group 1 causes and injuries, the decrease in YLLs was accompanied by a decrease in YLDs, albeit at a slower pace. The exceptions were neonatal encephalopathy, haemolytic disease and other neonatal jaundice, leishmaniasis, meningitis, hepatitis, and sexually transmitted diseases with increasing YLD rates between 1990 and 2015. A few Group 1 disorders and injuries had a faster decrease in YLDs than in YLLs: intestinal infections, obstructed labour, and neonatal sepsis. The only NCDs with a faster decrease in YLDs compared with YLLs were epilepsy and cervical cancer. Another small number of NCDs saw an increase in YLDs and YLLs, including drug use disorders, diabetes, and Parkinson's disease. NCDs with decreasing YLLs but increasing YLDs included

cancers of the prostate, testis, uterus, kidney, colorectum, and pancreas, melanoma, and congenital disorders. Some cancers (stomach and Hodgkin's lymphoma), rheumatic heart disease, asthma, chronic obstructive pulmonary disease, acute glomerulonephritis, peptic ulcer disease, gastritis, hernia, and gallbladder disease had decreasing YLLs and YLDs, but with a faster decrease in YLLs than in YLDs. The rate of change in YLDs for the main drivers of non-fatal health loss, musculoskeletal disorders, and mental and substance use disorders was small.

Global distribution of disability weights across individuals

Figure 5 shows the global distribution of individuals from our comorbidity microsimulation by six categories of disability, age, and sex for the highest and lowest SDI quintile. The six categories of disability are no disability, very mild disability (disability weight less than or equal to 0.01), mild disability (from 0.01 to 0.05 inclusive), moderate (from 0.05 to 0.1 inclusive), severe (from 0.1 to 0.3 inclusive), and profound (greater than 0.3). In 2015, most of the world's population experienced mild or greater disability. Having no disability at all was most common in children. After age 25 years, the proportion of the population having no disability became progressively smaller, and by age 55 years in low SDI countries and age 75 years in high SDI countries, nearly everyone had some form of disability. Very mild to moderate disability (ie, individuals with a disability weight of 0.1 or less) was common in childhood and young adults, but was replaced by more severe disability with increasing age. The patterns were similar for both sexes, apart from a much larger amount of disability in women older than 80 years in the top SDI quintiles, reflecting the much higher average age of women in this age category. For policy considerations around the age of retirement in ageing populations, it is noteworthy that from age 60 years onwards more than half of the population had severe or worse disability. The extent to which this loss of health limits or precludes the ability to work depends on the nature of the impairments and the type of employment in older workers with that level of disability.

Expected changes in disease profile with higher Socio-demographic Index

Figure 6 depicts changes in patterns of disability by level of SDI for age-standardised and all-age YLD rates per 100 000. Age-standardised YLD rates gradually decreased with increasing SDI in both sexes (figure 6A). The cause composition of YLDs somewhat varied across levels of SDI; these differences were largely derived from absolute levels of disability due to communicable causes and nutritional deficiencies, and to a lesser extent, maternal and neonatal disorders. Age-standardised YLD rates due to NCDs and injuries were similar at all SDI levels.

Across levels of SDI, mental and substance use disorders, musculoskeletal disorders, and other NCDs were consistently among the leading causes of

	Prevalence (thousands)		Percentage change (%)
	2005	2015	
(Continued from previous page)			
Other sense organ diseases	214 761 (207 401–222 622)	266 346 (257 047–276 383)	24.0 (23.1 to 24.9)*
Schistosomiasis	329 773 (297 093–367 878)	252 340 (211 032–321 081)	–23.5 (–32.5 to –4.4)*
G6PD deficiency	231 109 (205 067–259 769)	247 074 (209 307–286 713)	6.9 (1.4 to 11.9)*
Dermatitis	215 260 (199 590–230 536)	245 291 (227 283–262 752)	14.0 (13.2 to 14.8)*
Osteoarthritis	178 665 (173 558–184 053)	237 369 (230 336–244 648)	32.9 (31.9 to 33.8)*
Major depressive disorder	183 434 (163 947–206 420)	216 047 (192 863–243 319)	17.8 (16.6 to 19.0)*
Scabies	191 482 (166 101–223 739)	204 152 (177 534–237 466)	6.6 (4.0 to 9.5)*
Viral skin diseases	161 167 (152 218–170 651)	174 843 (165 156–185 072)	8.5 (8.0 to 9.0)*
Chronic obstructive pulmonary disease	149 115 (137 380–160 739)	174 483 (160 205–188 952)	17.0 (15.1 to 19.0)*
Genital prolapse	137 383 (121 623–154 875)	161 679 (142 335–182 566)	17.7 (15.2 to 20.0)*
Gastritis and duodenitis	135 993 (134 420–137 351)	157 060 (154 055–160 141)	15.5 (12.9 to 17.9)*
Peripheral vascular disease	115 109 (101 439–131 405)	154 651 (136 318–176 211)	34.4 (33.4 to 35.2)*
Uterine fibroids	126 797 (120 900–133 050)	151 115 (144 147–158 477)	19.2 (18.8 to 19.5)*
Hepatitis C	120 457 (108 080–133 129)	142 123 (126 978–157 045)	18.0 (16.7 to 19.2)*
Other mental and substance use disorders	107 895 (107 213–108 449)	128 178 (127 512–128 877)	18.8 (17.9 to 19.7)*
Iodine deficiency	103 701 (93 441–116 438)	110 920 (100 337–125 253)	7.0 (4.7 to 9.4)*
Ischaemic heart disease	87 511 (80 133–96 170)	110 193 (100 332–121 427)	25.9 (24.6 to 27.2)*
Benign prostatic hyperplasia	80 684 (70 338–90 853)	104 625 (90 730–118 244)	29.7 (27.5 to 32.0)*
Dysthymia	86 812 (74 974–99 103)	104 106 (90 398–118 969)	19.9 (18.4 to 21.5)*
Chronic kidney disease due to diabetes	79 184 (68 481–90 737)	100 824 (86 923–115 652)	27.3 (24.9 to 29.9)*
Chronic kidney disease due to other causes	74 917 (64 012–86 576)	94 553 (81 142–109 371)	26.2 (24.2 to 28.3)*
Idiopathic developmental intellectual disability	82 996 (47 588–117 109)	92 074 (52 280–130 411)	10.9 (9.5 to 11.9)*
Other cardiovascular and circulatory diseases	72 772 (68 090–77 763)	90 348 (84 711–96 659)	24.2 (22.7 to 25.5)*
Otitis media	83 022 (73 657–93 266)	86 393 (76 374–97 104)	4.1 (2.7% to 5.5)*
Psoriasis	67 753 (65 107–70 298)	79 700 (76 691–82 804)	17.6 (17.0 to 18.3)*
Other haemoglobinopathies and haemolytic anaemias	73 045 (72 600–73 470)	74 385 (73 898–74 862)	1.8 (0.9 to 2.7%)

Data in parentheses are 95% UIs. *Percentage changes that are statistically significant.

Table 2: Global prevalence of longer duration (more than 3 months) sequelae in 2005 and 2015 for all ages and both sexes combined, with percentage change between 2005 and 2015 for Level 4 causes with prevalence greater than 1%

age-standardised YLD rates. Anaemia led to generally higher rates of age-standardised YLDs in women than in men across levels of SDI, but the largest imbalance occurred at SDI levels between 0·10 and 0·50. Disability from injuries exacted a larger burden for men than for women, particularly at lower levels of SDI.

Without adjustments for population age structure (figure 6B), the effect of ageing populations and causes of disability that disproportionately affect older individuals become prominent. At all levels of SDI, total YLDs per 100 000 did not notably differ by sex; instead, the cause composition of disability showed greater differences. Below an SDI score of 0·25, communicable causes accounted for 30–45% of total disability, primarily due to nutritional deficiencies, malaria, and neglected tropical diseases. YLDs per 100 000 due to musculoskeletal disorders, particularly lower back and neck pain and other musculoskeletal disorders, increased substantially from low to high SDI, with a more pronounced increase beginning at an SDI score of 0·6. This rise was particularly evident in women.

Trends in age-standardised YLDs per capita

Globally, age-standardised YLDs per capita (an indicator of overall disability experienced per person in a given place) moderately decreased for both sexes between 1990 and 2015 (results appendix pp 714–16). Age-standardised YLDs per capita were consistently higher for women than for men at the global level. For both sexes, YLDs per capita were generally higher for lower levels of SDI. YLDs per capita were noticeably larger for low SDI and low-middle SDI groups than for other SDI levels (ie, high SDI, high-middle SDI, and middle SDI), which were more similar to each other.

Leading causes of YLDs and deviations from expected levels based on Socio-demographic Index

Clear, though varied, patterns emerged across and within GBD regions in comparison of observed levels of YLDs due to leading causes of disability with levels expected on the basis of SDI. Figure 7 displays ratios of observed and expected YLDs for the leading ten causes at level 3 of the GBD hierarchy in 2015, colour coded by the magnitude of differences between observed and expected YLDs.

Globally, lower back and neck pain was the leading cause of disability in 2015. Two mental disorders, major depressive and anxiety disorders, were the third and ninth leading causes of global disability, and diabetes was the sixth leading driver of disability. Iron-deficiency anaemia was the only Group 1 cause among the leading ten causes for global YLDs (ranked fourth). Sensory disorders ranked second and skin diseases ranked fifth. Lower back and neck pain was the leading global cause of disability in 2015 in most countries. The leading cause of disability in 2015 was iron-deficiency anaemia in 27 countries; HIV/AIDS in all six southern sub-Saharan

African countries; depression in five eastern sub-Saharan Africa countries; other neglected tropical diseases in Angola, Democratic Republic of the Congo, and Gabon; sense organ disorders in Comoros and Myanmar; diabetes in Fiji and Marshall Islands; war in Lebanon and Syria; and onchocerciasis in Liberia.

Regional, country, territory, and selected subnational results

Lower back and neck pain was the leading cause of disability in all high-income countries in 2015. However, ratios of observed to expected YLDs from lower back and neck pain ranged from 0·60 for Singapore to more than 1·59 in Norway. Most high-income countries experienced higher than expected levels of disability due to depressive disorders. The USA and Australia were the only two high-income countries where drug use disorders were a top ten cause of disability, and observed levels of YLDs were much higher than expected. South Korea's ratio of observed-to-expected levels of YLDs due to diabetes exceeded 1·30, whereas Japan's diabetes-related disability was lower than expected on the basis of SDI. In the UK, observed disability due to asthma was well above expected levels.

In 2015, lower back and neck pain was the leading cause of YLDs for all but two countries in Latin America and the Caribbean; Haiti and Venezuela were the exceptions, with iron-deficiency anaemia as the leading causes of disability for both countries. Disability due to diabetes surpassed expected levels by at least a factor of two for six countries and territories: Antigua, Barbados, Dominica, Puerto Rico, Trinidad and Tobago, and the Virgin Islands. Peru, however, had a ratio of less than 0·6 for observed and expected YLDs for diabetes.

In 2015, lower back and neck pain was the leading cause of YLDs for 24 out of 28 countries and territories of southeast Asia, east Asia, and Oceania. Other leading causes of disability were sensory disorders in Myanmar, diabetes in Fiji and the Marshall Islands, and iron-deficiency anaemia in Papua New Guinea. Although lower back and neck pain was the primary cause of disability, many countries had far lower levels of this than expected given their SDI, including Thailand (0·73), Indonesia (0·76), and Malaysia (0·75). Observed YLDs due to depressive disorders were often lower than expected, with 22 countries recording ratios below 0·80. Conversely, numerous geographies recorded YLD ratios exceeding 2·00 for diabetes (eg, 2·29 in Taiwan).

Beyond lower back and neck pain, which was the leading cause of YLDs for three of five countries in south Asia in 2015 (with India and Pakistan being the exception), a mixture of causes accounted for the region's main causes of disability; this heterogeneity probably reflects the diversity of countries in the region and their places along the development spectrum. Iron-deficiency anaemia was the first leading cause and lower back and neck pain was the second leading cause of YLDs in both India and Pakistan, whereas sensory disorders, other musculoskeletal disorders, and iron-deficiency anaemia ranked second for

	Prevalence (thousands)			YLDs (thousands)		
	2015	Percentage change in counts between 2005 and 2015	Percentage change in ASR between 2005 and 2015	2015	Percentage change in counts between 2005 and 2015	Percentage change in ASR between 2005 and 2015
All causes	792 004.7 (588 538.7–1 019 955.2)	15.1 (14.5 to 15.6)*	-2.1 (-2.5 to -1.7)*
Communicable, maternal, neonatal, and nutritional diseases	4 133 822.3 (4 108 132.5 to 4 161 262.6)	8.0 (7.3 to 8.7)*	-4.2 (-4.8 to -3.6)*	112 501.9 (80 347.4–156 270.9)	-0.5 (-2.4 to 2.5)	-9.7 (-11.4 to -7.0)*
HIV/AIDS and tuberculosis	46 005.2 (44 985.8 to 47 146.6)	18.6 (15.2 to 22.1)*	2.9 (-0.0 to 5.9)	6702.3 (4798.6–8863.5)	2.1 (-3.5 to 7.2)	-12.2 (-17.0 to -7.8)*
Tuberculosis	8861.2 (8076.3 to 9707.2)	9.2 (6.3 to 12.2)*	-7.2 (-9.5 to -4.7)*	2712.4 (1829.9–3744.0)	9.4 (6.4 to 12.5)*	-6.8 (-9.2 to -4.2)*
HIV/AIDS	37 277.5 (36 279.9 to 38 339.1)	21.1 (16.9 to 25.5)*	5.6 (2.0 to 9.6)*	3989.9 (2891.2–5159.5)	-2.4 (-10.5 to 5.5)	-15.6 (-23.0 to -8.5)*
HIV/AIDS—tuberculosis	1258.0 (1142.5 to 1386.9)	-17.9 (-20.7 to -14.9)*	-28.3 (-30.7 to -25.7)*	471.9 (316.0–645.6)	-17.8 (-20.7 to -14.7)*	-28.1 (-30.6 to -25.5)*
HIV/AIDS resulting in other diseases	37 543.3 (36 350.0 to 39 028.4)	23.6 (20.5 to 26.7)*	7.9 (5.1 to 10.6)*	3518.0 (2 555.4–4 558.9)	0.2 (-9.3 to 9.3)	-13.6 (-22.1 to -5.2)*
HIV aggregate	21 149.0 (20 260.3 to 22 292.3)	-19.5 (-22.5 to -16.3)*	-29.7 (-32.3 to -26.9)*	1543.6 (1 107.2–2 025.2)	-20.4 (-23.4 to -17.2)*	-30.4 (-33.1 to -27.6)*
AIDS aggregate	16 394.3 (15 747.0 to 17 154.2)	299.5 (251.4 to 355.5)*	232.9 (193.5 to 279.0)*	1974.4 (1432.0–2559.9)	25.5 (0.9 to 56.1)*	6.1 (-14.6 to 32.4)
Diarrhoea, lower respiratory infections, and other common infectious diseases	402 320.4 (393 746.5 to 408 994.2)	8.0 (7.2 to 8.8)*	-2.9 (-3.7 to -2.3)*	14 865.0 (10 397.0–20 283.4)	6.2 (5.4 to 7.1)*	-4.0 (-4.7 to -3.3)*
Diarrhoeal diseases	35 820.6 (34 342.1 to 37 537.8)	6.4 (5.5 to 7.3)*	-3.3 (-4.1 to -2.5)*	5731.7 (3 943.3–7 890.5)	6.4 (5.4 to 7.5)*	-3.1 (-4.0 to -2.2)*
Diarrhoea episodes aggregate	35 816.3 (34 337.5 to 37 534.1)	6.4 (5.5 to 7.3)*	-3.3 (-4.1 to -2.5)*	5730.4 (3 942.2–7 889.0)	6.4 (5.4 to 7.5)*	-3.1 (-4.0 to -2.2)*
Guillain-Barré syndrome due to diarrhoeal diseases	4.2 (3.2 to 5.5)	16.9 (14.7 to 19.5)*	-0.2 (-0.9 to 0.4)	1.3 (0.8–1.9)	16.9 (14.7 to 19.5)*	-0.2 (-0.9 to 0.4)
Intestinal infectious diseases	1666.8 (1594.4 to 1 732.8)	-23.3 (-25.6 to -20.6)*	-28.2 (-30.4 to -25.8)*	220.3 (149.3–306.5)	-18.6 (-22.8 to -13.8)*	-24.2 (-28.1 to -19.8)*
Typhoid fever	1446.7 (1256.2 to 1 648.1)	-18.3 (-20.8 to -15.6)*	-23.9 (-26.3 to -21.3)*	191.0 (129.5–267.8)	-17.7 (-22.1 to -12.6)*	-23.2 (-27.4 to -18.5)*
Typhoid fever episodes aggregate	1197.9 (1041.3 to 1366.6)	-18.3 (-21.7 to -14.8)*	-23.9 (-27.1 to -20.5)*	113.3 (75.8–162.5)	-18.1 (-22.6 to -13.4)*	-23.7 (-27.8 to -19.2)*
Complications of typhoid fever aggregate	248.8 (210.8 to 292.2)	-18.2 (-28.6 to -5.4)*	-23.8 (-33.5 to -12.1)*	77.8 (51.6–110.8)	-17.0 (-28.5 to -3.0)*	-22.6 (-33.1 to -9.9)*
Paratyphoid fever	529.3 (431.3 to 646.3)	-17.8 (-22.6 to -12.7)*	-23.8 (-28.2 to -19.2)*	27.5 (17.5–40.7)	-17.5 (-24.7 to -9.2)*	-23.5 (-30.1 to -16.1)*
Paratyphoid fever episodes aggregate	502.2 (409.6 to 612.7)	-17.8 (-22.6 to -12.4)*	-23.8 (-28.2 to -19.0)*	24.4 (15.5–36.4)	-17.5 (-24.9 to -8.7)*	-23.5 (-30.7 to -15.5)*
Intestinal perforation due to paratyphoid	27.1 (20.9 to 34.5)	-17.6 (-33.0 to 1.4)	-23.7 (-37.8 to -6.1)*	3.1 (1.9–4.6)	-17.5 (-32.8 to 1.6)	-23.6 (-37.7 to -5.9)*
Other intestinal infectious diseases	1.8 (0.6–4.1)	-67.0 (-78.8 to -45.8)*	-69.3 (-80.2 to -50.1)*
Lower respiratory infections	8986.6 (8545.2 to 9393.7)	5.1 (4.0 to 6.2)*	-8.2 (-9.1 to -7.3)*	540.4 (365.3–760.3)	4.9 (3.5 to 6.3)*	-8.1 (-9.2 to -7.0)*
Lower respiratory infection episodes aggregate	8982.2 (8539.8 to 9387.9)	5.1 (4.0 to 6.2)*	-8.2 (-9.1 to -7.3)*	539.1 (364.5–759.5)	4.9 (3.5 to 6.3)*	-8.1 (-9.2 to -7.0)*
Guillain-Barré syndrome due to lower respiratory infections	4.4 (2.5 to 6.9)	16.9 (14.7 to 19.5)*	-0.2 (-0.9 to 0.4)	1.3 (0.7–2.2)	16.9 (14.7 to 19.5)*	-0.2 (-0.9 to 0.4)
Upper respiratory infections	233 470.3 (208 009.7 to 259 076.8)	10.2 (8.9 to 11.5)*	-1.4 (-2.0 to -0.8)*	2738.4 (1538.6–4644.3)	10.1 (8.9 to 11.5)*	-1.3 (-1.9 to -0.6)*
Upper respiratory infection episodes aggregate	233 458.2 (207 997.9 to 259 065.2)	10.2 (8.9 to 11.5)*	-1.4 (-2.0 to -0.8)*	2734.8 (1 533.9–4 641.4)	10.1 (8.9 to 11.5)*	-1.3 (-1.9 to -0.6)*
Guillain-Barré syndrome due to upper respiratory infections	12.1 (9.6 to 15.2)	16.9 (14.7 to 19.5)*	-0.2 (-0.9 to 0.4)	3.6 (2.2–5.4)	17.0 (14.7 to 19.5)*	-0.2 (-0.9 to 0.4)

(Table 3 continues on next page)

	Prevalence (thousands)			YLDs (thousands)		
	2015	Percentage change in counts between 2005 and 2015	Percentage change in ASR between 2005 and 2015	2015	Percentage change in counts between 2005 and 2015	Percentage change in ASR between 2005 and 2015
(Continued from previous page)						
Otitis media	112 089.1 (100 504.6 to 123 655.6)	5.2 (4.1 to 6.5)*	-4.4 (-5.5 to -3.4)*	3321.5 (2 114.2-4 906.3)	3.3 (0.6 to 6.0)*	-5.6 (-8.0 to -3.1)*
Acute otitis media	25 630.8 (21 044.2 to 31 570.6)	9.4 (7.5 to 11.2)*	1.3 (-0.4 to 3.0)	334.9 (169.4-623.7)	9.4 (7.5 to 11.4)*	1.4 (-0.3 to 3.1)
Chronic otitis media aggregate	86 458.3 (76 403.0 to 97 133.7)	4.1 (2.7 to 5.5)*	-6.0 (-7.2 to -4.8)*	2986.6 (1904.2-4 371.1)	2.7 (-0.2 to 5.6)	-6.3 (-8.9 to -3.6)*
Meningitis	8733.5 (8320.6 to 9107.1)	16.9 (13.4 to 20.0)*	2.5 (-0.6 to 5.1)	1516.9 (1076.1-1986.0)	15.0 (13.7 to 16.5)*	1.9 (0.7 to 3.0)*
Pneumococcal meningitis	7297.2 (6228.5 to 8635.3)	22.2 (19.7 to 25.1)*	6.9 (4.7 to 9.6)*	728.8 (513.1-959.9)	17.6 (15.5 to 19.7)*	3.9 (2.0 to 5.6)*
Acute pneumococcal meningitis	40.3 (33.5 to 48.4)	25.3 (17.4 to 33.8)*	12.1 (5.2 to 19.5)*	5.3 (3.4-7.8)	25.1 (15.1 to 37.1)*	11.9 (3.2 to 22.3)*
Complications of pneumococcal meningitis aggregate	7256.9 (6191.3 to 8591.7)	22.2 (19.7 to 25.1)*	6.9 (4.7 to 9.6)*	723.5 (509.2-952.4)	17.6 (15.4 to 19.6)*	3.8 (2.0 to 5.6)*
H influenzae type b meningitis	2455.6 (1966.7 to 3032.6)	-8.3 (-12.2 to -3.9)*	-18.0 (-21.5 to -13.9)*	302.1 (209.4-407.1)	6.9 (3.4 to 10.6)*	-3.3 (-6.6 to 0.0)
Acute H influenzae type b meningitis	22.0 (17.2 to 28.7)	-19.2 (-25.3 to -11.9)*	-26.0 (-31.8 to -19.2)*	2.9 (1.8-4.5)	-18.7 (-26.1 to -10.8)*	-25.6 (-32.4 to -18.1)*
Complications of H influenzae type b meningitis aggregate	2433.5 (1947.4 to 3010.8)	-8.2 (-12.1 to -3.7)*	-17.9 (-21.5 to -13.7)*	299.1 (207.9-402.7)	7.3 (3.7 to 11.0)*	-3.0 (-6.3 to 0.3)
Meningococcal meningitis	1720.8 (1327.7 to 2 180.5)	22.0 (17.9 to 26.2)*	7.5 (3.8 to 11.3)*	161.6 (111.7-215.6)	19.7 (17.5 to 22.1)*	5.1 (3.2 to 7.4)*
Acute meningococcal meningitis	22.8 (18.9 to 27.7)	24.4 (14.4 to 34.9)*	12.7 (3.7 to 22.4)*	3.0 (1.8-4.5)	24.7 (10.9 to 39.4)*	12.9 (0.5 to 25.9)*
Complications of meningococcal meningitis aggregate	1698.0 (1305.2 to 2155.2)	22.0 (17.9 to 26.2)*	7.4 (3.8 to 11.2)*	158.6 (109.4-211.9)	19.6 (17.4 to 22.0)*	5.0 (3.1 to 7.3)*
Other meningitis	3015.2 (2438.5 to 3618.0)	19.0 (16.6 to 21.6)*	3.5 (1.4 to 5.9)*	324.5 (225.1-428.1)	15.3 (13.3 to 17.4)*	0.9 (-0.8 to 2.7)
Other meningitis episodes aggregate	134.5 (120.0 to 151.7)	16.3 (10.2 to 22.5)*	5.7 (0.1 to 11.4)*	17.8 (11.5-25.3)	16.6 (8.6 to 24.0)*	5.9 (-1.1 to 12.5)
Complications of other meningitis aggregate	2880.4 (2300.7 to 3474.9)	19.1 (16.5 to 21.9)*	3.4 (1.2 to 5.8)*	306.7 (213.3-406.6)	15.2 (13.2 to 17.5)*	0.6 (-1.1 to 2.6)
Encephalitis	4315.8 (3145.8 to 5875.6)	4.8 (2.8 to 7.4)*	-9.3 (-11.3 to -6.7)*	457.0 (326.7-594.9)	6.7 (4.6 to 9.0)*	-6.3 (-8.2 to -4.3)*
Acute encephalitis	81.8 (74.8 to 89.5)	7.1 (5.2 to 9.0)*	-4.1 (-5.7 to -2.4)*	10.8 (7.1-15.5)	7.5 (3.8 to 11.1)*	-3.7 (-6.9 to -0.5)*
Complications of encephalitis aggregate	4233.6 (3069.6 to 5787.5)	4.8 (2.7 to 7.3)*	-9.4 (-11.4 to -6.8)*	446.2 (318.8-581.4)	6.7 (4.6 to 9.0)*	-6.4 (-8.3 to -4.4)*
Diphtheria	0.6 (0.3 to 1.2)	-59.6 (-81.7 to -12.7)*	-62.8 (-83.1 to -19.5)*	0.0 (0.0-0.1)	-59.7 (-81.7 to -12.7)*	-62.8 (-83.1 to -19.5)*
Whooping cough	2232.6 (1725.9 to 2800.7)	-27.4 (-29.5 to -25.2)*	-32.4 (-34.3 to -30.3)*	110.3 (65.6-167.0)	-32.3 (-29.6 to -24.8)*	-32.3 (-34.5 to -30.0)*
Tetanus	209.3 (204.8 to 214.7)	8.4 (6.1 to 9.8)*	-2.7 (-5.0 to -1.4)*	9.1 (6.7-12.2)	-8.9 (-14.5 to -3.7)*	-18.1 (-23.3 to -13.3)*
Severe tetanus	8.6 (6.5 to 13.0)	-46.3 (-54.0 to -36.0)*	-52.1 (-59.1 to -42.9)*	1.1 (0.7-1.9)	-45.9 (-53.8 to -35.7)*	-51.8 (-58.9 to -42.6)*
Complications of tetanus aggregate	200.7 (196.7 to 204.5)	13.3 (12.8 to 13.9)*	1.8 (1.4 to 2.3)*	8.0 (5.8-10.6)	1.0 (-3.7 to 5.4)	-8.8 (-13.0 to -4.9)*
Measles	127.4 (56.8 to 252.2)	-70.2 (-73.6 to -66.0)*	-72.3 (-75.4 to -68.4)*	11.5 (4.3-25.3)	-70.0 (-73.6 to -65.4)*	-72.0 (-75.4 to -67.7)*
Varicella and herpes zoster	5907.7 (5489.4 to 6344.6)	15.2 (14.1 to 16.5)*	-0.4 (-0.8 to -0.0)*	207.8 (127.9-316.5)	18.5 (16.5 to 20.8)*	-0.4 (-1.4 to 0.7)

(Table 3 continues on next page)

	Prevalence (thousands)			YLDs (thousands)		
	2015	Percentage change in counts between 2005 and 2015	Percentage change in ASR between 2005 and 2015	2015	Percentage change in counts between 2005 and 2015	Percentage change in ASR between 2005 and 2015
(Continued from previous page)						
Neglected tropical diseases and malaria	1 900 062.6 (1 863 148.8 to 1 940 058.5)	-0.4 (-2.5 to 1.9)	-10.7 (-12.7 to -8.7)*	20 763.1 (13 382.4-32 174.5)	-4.7 (-13.2 to 8.9)	-14.5 (-22.2 to -2.4)*
Malaria	295 717.3 (257 568.4 to 338 449.0)	29.9 (22.5 to 37.2)*	20.0 (12.9 to 27.2)*	3358.2 (2356.9-4703.9)	16.1 (12.8 to 19.3)*	8.5 (5.4 to 11.4)*
Asymptomatic malaria parasitaemia (PfPR)	206 147.2 (168 670.7 to 246 888.7)	37.3 (27.0 to 48.1)*	26.4 (16.6 to 36.8)*	0.0 (0.0-0.0)	0.0 (0.0 to 0.0)	0.0 (0.0 to 0.0)
Malaria episodes aggregate	16 756.8 (10 341.8 to 25 965.5)	-15.9 (-23.6 to -8.4)*	-21.1 (-28.4 to -14.1)*	346.6 (174.4-598.4)	-15.8 (-23.6 to -8.2)*	-20.9 (-28.2 to -13.7)*
Complications of malaria aggregate	876.9 (798.1 to 957.9)	28.4 (26.5 to 30.5)*	18.4 (16.5 to 20.3)*	274.9 (209.7-344.8)	24.7 (20.5 to 29.1)*	15.0 (11.2 to 18.9)*
Anaemia due to malaria parasitaemia (PfPR) aggregate	71 936.4 (70 137.5 to 73 504.2)	26.3 (24.4 to 28.0)*	17.0 (15.3 to 18.6)*	2736.7 (1848.0-3918.8)	21.1 (18.1 to 24.3)*	13.1 (10.5 to 16.1)*
Chagas disease	6653.6 (5750.5 to 7575.6)	-7.1 (-9.7 to -4.1)*	-22.1 (-24.2 to -19.7)*	63.1 (42.1-90.8)	-1.8 (-5.1 to 1.7)	-21.0 (-23.6 to -18.3)*
Chagas disease episodes aggregate	5634.7 (4850.5 to 6414.1)	-7.8 (-10.3 to -4.9)*	-22.2 (-24.3 to -19.8)*	0.0 (0.0-0.1)	-18.7 (-22.9 to -14.4)*	-27.6 (-31.1 to -24.1)*
Complications of Chagas disease aggregate	666.0 (525.5 to 820.6)	-5.2 (-7.9 to -1.9)*	-22.0 (-24.3 to -19.6)*	44.8 (30.0-62.8)	-2.5 (-6.0 to 1.2)	-21.0 (-23.6 to -18.3)*
Heart failure due to Chagas disease aggregate	353.0 (221.5 to 503.5)	0.3 (-2.9 to 4.0)	-20.9 (-23.4 to -18.2)*	18.2 (9.7-29.4)	0.2 (-3.6 to 4.4)	-20.9 (-23.8 to -17.8)*
Leishmaniasis	3859.3 (3438.4 to 4570.4)	27.3 (5.9 to 55.0)*	11.4 (-7.0 to 35.1)	45.8 (22.8-86.7)	25.5 (21.9 to 28.9)*	10.5 (7.5 to 13.5)*
Visceral leishmaniasis	60.8 (57.5 to 64.7)	10.6 (9.7 to 11.4)*	2.3 (1.6 to 3.0)*	4.3 (2.9-6.1)	10.8 (2.1 to 20.7)*	2.6 (-5.4 to 11.5)
Cutaneous and mucocutaneous leishmaniasis	3895.9 (3 324.6 to 4 767.5)	27.0 (23.7 to 30.1)*	11.0 (8.3 to 13.8)*	41.5 (19.4-80.8)	27.3 (23.6 to 30.5)*	11.4 (8.3 to 14.5)*
African trypanosomiasis	10.7 (6.0 to 17.0)	-68.7 (-70.4 to -67.3)*	-72.3 (-73.8 to -71.0)*	3.0 (1.4-5.3)	-67.4 (-72.6 to -62.1)*	-71.0 (-75.5 to -66.6)*
Schistosomiasis	252 339.5 (211 032.5 to 321 081.3)	-23.5 (-32.5 to -4.4)*	-30.9 (-39.0 to -13.7)*	2472.6 (1275.0-4521.2)	-21.8 (-29.3 to -4.0)*	-29.1 (-36.0 to -13.1)*
Schistosomiasis episodes aggregate	130 285.7 (114 433.9 to 156 900.0)	-26.3 (-34.5 to -13.6)*	-33.1 (-40.5 to -21.3)*	740.7 (292.3-1584.0)	-27.3 (-35.2 to -14.1)*	-34.0 (-41.0 to -22.0)*
Complications of schistosomiasis aggregate	105 621.9 (76 641.0 to 160 339.4)	-19.6 (-32.6 to 14.1)	-27.7 (-39.4 to 3.0)	1188.8 (535.9-2 463.3)	-18.0 (-30.5 to 15.6)	-26.3 (-37.5 to 4.3)
Anaemia due to schistosomiasis aggregate	16 331.3 (16 075.8 to 16 655.1)	-19.6 (-21.1 to -17.7)*	-27.0 (-28.3 to -25.3)*	543.1 (365.1-781.5)	-21.5 (-24.0 to -18.6)*	-27.8 (-30.0 to -25.1)*
Cysticercosis	1931.0 (1597.8 to 2312.0)	-6.2 (-10.2 to -2.5)*	-20.8 (-24.3 to -17.6)*	286.7 (194.2-392.8)	-16.3 (-21.3 to -11.8)*	-29.2 (-33.3 to -25.3)*
Cystic echinococcosis	1383.0 (1265.9 to 1498.6)	24.7 (22.6 to 27.0)*	6.7 (5.0 to 8.5)*	126.8 (86.7-174.6)	24.3 (21.1 to 27.5)*	6.6 (4.1 to 9.2)*
Lymphatic filariasis	38 464.1 (31 328.2 to 46 783.0)	-44.9 (-49.9 to -39.9)*	-51.6 (-56.0 to -47.2)*	2075.0 (1120.6-3311.2)	-16.2 (-32.1 to -3.9)*	-27.7 (-41.5 to -17.1)*
Prevalence of detectable microfilaria due to lymphatic filariasis	19 707.9 (17 173.8 to 22 270.5)	-58.8 (-61.8 to -55.3)*	-63.5 (-66.1 to -60.3)*	0.0 (0.0-0.0)	0.0 (0.0 to 0.0)	0.0 (0.0 to 0.0)
Complications of lymphatic filariasis aggregate	18 756.3 (12 420.1 to 26 397.1)	-14.8 (-29.9 to -3.5)*	-26.7 (-40.1 to -16.7)*	2075.0 (1120.6-3311.2)	-16.2 (-32.1 to -3.9)*	-27.7 (-41.5 to -17.1)*
Onchocerciasis	15 531.5 (11 963.5 to 19 993.8)	-29.1 (-39.5 to -18.8)*	-36.8 (-46.0 to -27.5)*	1135.7 (546.2-2005.4)	-21.2 (-38.5 to -4.8)*	-31.2 (-46.7 to -16.2)*
Asymptomatic onchocerciasis	2280.4 (1525.0 to 3173.2)	-62.4 (-66.1 to -59.6)*	-66.2 (-69.5 to -63.7)*	0.0 (0.0-0.0)	0.0 (0.0 to 0.0)	0.0 (0.0 to 0.0)

(Table 3 continues on next page)

(Continued from previous page)

	Prevalence (thousands)			YLDs (thousands)		
	2015	Percentage change in counts between 2005 and 2015	Percentage change in ASR between 2005 and 2015	2015	Percentage change in counts between 2005 and 2015	Percentage change in ASR between 2005 and 2015
Skin disease due to onchocerciasis aggregate	12 223.9 (8 474.7 to 16 585.2)	-17.4 (-31.0 to -2.7)*	-26.5 (-38.6 to -13.1)*	1055.6 (467.4-1894.7)	-22.2 (-41.8 to -3.2)*	-31.6 (-49.0 to -14.8)*
Vision loss due to onchocerciasis aggregate	1025.4 (724.8 to 1452.1)	-2.3 (-15.0 to 11.8)	-22.3 (-32.3 to -11.0)*	80.1 (51.3-117.5)	-6.1 (-16.9 to 6.0)	-25.4 (-33.9 to -15.9)*
Trachoma	3557.1 (2940.5 to 4321.8)	4.6 (0.4 to 9.2)*	-19.3 (-22.9 to -15.2)*	279.2 (192.8-396.0)	-1.2 (-5.6 to 3.0)	-23.9 (-27.5 to -20.3)*
Dengue	4730.0 (2654.1 to 10 254.2)	143.6 (-0.3 to 564.7)	119.7 (-10.1 to 498.7)	764.1 (346.8-1 744.2)	140.8 (-0.1 to 558.4)	117.7 (-9.8 to 494.3)
Dengue episodes aggregate	1409.2 (937.6 to 2943.8)	134.1 (-3.8 to 521.2)	110.8 (-13.2 to 459.0)	82.1 (45.0-183.9)	141.4 (0.4 to 549.2)*	117.9 (-9.4 to 486.7)
Post-dengue chronic fatigue syndrome	3324.6 (1600.3 to 7347.9)	143.7 (-0.3 to 564.7)	119.8 (-10.1 to 498.7)	682.0 (295.2-1 608.9)	140.8 (-0.2 to 558.0)	117.6 (-9.8 to 494.1)
Yellow fever	2.8 (0.8 to 7.7)	-25.7 (-31.6 to -19.2)*	-31.6 (-36.9 to -25.8)*	0.1 (0.0-0.3)	-25.7 (-31.6 to -19.2)*	-31.6 (-36.9 to -25.8)*
Rabies	0.7 (0.6 to 0.8)	-43.4 (-51.5 to -33.8)*	-50.8 (-57.6 to -42.6)*	0.1 (0.1-0.1)	-43.4 (-51.5 to -33.8)*	-50.8 (-57.6 to -42.6)*
Intestinal nematode infections	1 447 209.3 (1 414 995.0 to 1 481 332.3)	-2.5 (-5.1 to 0.2)	-12.8 (-15.2 to -10.4)*	3173.8 (1861.3-5090.8)	-22.8 (-27.3 to -17.3)*	-30.4 (-34.5 to -25.4)*
Ascariasis	761 893.8 (682 557.5 to 861 031.5)	-10.8 (-22.5 to 2.5)	-20.3 (-30.7 to -8.2)*	871.0 (482.5-1 464.3)	-37.7 (-45.3 to -29.4)*	-43.9 (-50.8 to -36.4)*
Complications of ascariasis aggregate	47 626.4 (44 087.5 to 51 711.9)	-35.6 (-41.9 to -28.4)*	-42.1 (-47.8 to -35.6)*	871.0 (482.5-1 464.3)	-37.7 (-45.3 to -29.4)*	-43.9 (-50.8 to -36.4)*
Asymptomatic ascariasis	714 230.7 (634 201.9 to 814 009.5)	-8.5 (-20.9 to 6.4)	-18.2 (-29.3 to -4.9)*	0.0 (0.0-0.0)	0.0 (0.0 to 0.0)	0.0 (0.0 to 0.0)
Trichuriasis	463 652.1 (426 621.2 to 502 939.4)	-2.1 (-12.0 to 9.0)	-12.2 (-21.1 to -2.1)*	544.1 (290.7-946.0)	-16.7 (-30.6 to 3.6)	-24.9 (-37.5 to -6.8)*
Complications of trichuriasis aggregate	27 129.6 (24 188.7 to 31 890.8)	-15.6 (-27.8 to 2.1)	-24.1 (-35.0 to -8.2)*	544.1 (290.7-946.0)	-16.7 (-30.6 to 3.6)	-24.9 (-37.5 to -6.8)*
Asymptomatic trichuriasis	436 525.4 (400 013.3 to 475 985.5)	-1.1 (-11.8 to 11.0)	-11.3 (-21.0 to -0.3)*	0.0 (0.0-0.0)	0.0 (0.0 to 0.0)	0.0 (0.0 to 0.0)
Hookworm disease	428 245.9 (394 486.0 to 468 292.4)	-7.3 (-16.9 to 3.5)	-17.2 (-25.8 to -7.5)*	1758.8 (1088.5-2754.9)	-14.6 (-19.9 to -9.4)*	-22.9 (-27.7 to -18.2)*
Complications of hookworm disease aggregate	51 012.9 (48 151.1 to 54 346.2)	-14.0 (-21.0 to -6.2)*	-23.2 (-29.5 to -16.3)*	1046.7 (572.8-1732.1)	-15.5 (-24.0 to -6.3)*	-24.4 (-32.1 to -16.3)*
Anaemia due to hookworm disease aggregate	24 220.2 (24 078.2 to 24 365.2)	-6.2 (-7.3 to -5.1)	-15.5 (-16.5 to -14.6)*	712.1 (476.3-1031.1)	-13.2 (-15.6 to -11.1)*	-20.5 (-22.6 to -18.7)*
Asymptomatic hookworm disease	352 948.6 (319 143.2 to 392 574.4)	-6.4 (-18.3 to 7.5)	-16.4 (-27.2 to -3.9)*	0.0 (0.0-0.0)	0.0 (0.0 to 0.0)	0.0 (0.0 to 0.0)
Food-borne trematodiasis	71 095.4 (67 365.5 to 75 246.9)	4.0 (1.5 to 6.6)*	-9.3 (-11.4 to -7.1)*	1686.5 (857.1-3066.8)	3.7 (-0.4 to 10.1)*	-10.0 (-13.4 to -5.2)*
Asymptomatic food-borne trematodiasis aggregate	56 131.7 (45 765.9 to 63 620.3)	3.8 (1.2 to 6.6)*	-9.3 (-11.6 to -6.9)*	0.0 (0.0-0.0)	0.0 (0.0 to 0.0)	0.0 (0.0 to 0.0)
Symptomatic food-borne trematodiasis aggregate	14 963.7 (8 632.7 to 25 256.6)	4.8 (0.5 to 11.4)*	-9.3 (-12.8 to -4.2)*	1686.5 (857.1-3066.8)	3.7 (-0.4 to 10.1)	-10.0 (-13.4 to -5.2)*
Leprosy	514.2 (487.0 to 545.7)	-0.1 (-0.5 to 0.3)	-19.7 (-20.2 to -19.3)*	31.0 (20.9-43.5)	0.6 (-1.5 to 2.9)	-19.2 (-20.8 to -17.4)*
Ebola	2.8 (1.2 to 5.2)	54 521.3 (46 386.2 to 68 346.9)*	49 817.9 (41 865.6 to 62 764.9)*	0.6 (0.2-1.1)	51 908.8 (42 739.0 to 66 884.3)*	47 453.0 (38 609.5 to 61 730.3)*
Other neglected tropical diseases	61 452.6 (60 842.8 to 62 054.9)	0.1 (-1.0 to 1.3)	-7.7 (-8.8 to -6.6)*	5261.1 (2558.5-12 199.0)	7.4 (-18.9 to 50.6)	-1.2 (-25.5 to 38.0)

(Table 3 continues on next page)

	Prevalence (thousands)			YLDs (thousands)		
	2015	Percentage change in counts between 2005 and 2015	Percentage change in ASR between 2005 and 2015	2015	Percentage change in counts between 2005 and 2015	Percentage change in ASR between 2005 and 2015
(Continued from previous page)						
Anaemia due to other neglected tropical diseases aggregate	61 452.6 (60 842.8 to 62 054.9)	0.1 (-1.0 to 1.3)	-7.7 (-8.8 to -6.6)*	2158.2 (1 446.3-3 073.1)	-6.0 (-8.2 to -3.8)*	-12.4 (-14.5 to -10.4)*
Maternal disorders	11 372.9 (10 686.6 to 11 894.1)	2.1 (-4.2 to 9.2)	-7.9 (-13.6 to -1.8)*	898.8 (642.8-1222.9)	-5.1 (-18.7 to 11.2)	-15.1 (-27.0 to -0.3)*
Maternal haemorrhage	2304.8 (2231.6 to 2 393.5)	1.6 (-3.0 to 6.5)	-7.8 (-12.0 to -3.4)*	84.7 (57.3-119.5)	0.1 (-13.5 to 17.2)	-9.1 (-21.3 to 6.6)
Maternal hemorrhage episodes aggregate	191.7 (127.8 to 277.5)	2.9 (-36.5 to 66.2)	-6.4 (-42.1 to 49.9)	31.3 (18.1-49.0)	3.3 (-30.6 to 54.9)	-6.0 (-36.3 to 41.1)*
Anaemia due to maternal haemorrhage aggregate	2113.4 (2082.8 to 2141.5)	1.5 (-0.9 to 3.9)	-7.9 (-10.1 to -5.8)*	53.5 (35.0-78.6)	-1.8 (-5.9 to 2.6)	-10.8 (-14.5 to -6.9)*
Maternal sepsis and other maternal infections	3140.8 (2658.0 to 3727.7)	3.7 (-15.2 to 27.8)	-7.8 (-24.2 to 13.1)	56.5 (29.5-102.4)	-1.0 (-42.3 to 70.4)	-9.7 (-47.1 to 55.2)
Maternal sepsis and other maternal infections aggregate	900.0 (486.7 to 1462.3)	-2.4 (-50.3 to 93.7)	-10.2 (-54.4 to 78.1)	45.3 (22.2-82.7)	-2.7 (-49.0 to 91.0)	-10.5 (-53.2 to 73.6)
Infertility due to puerperal sepsis	2242.1 (2 030.5 to 2483.0)	6.4 (4.4 to 8.9)*	-6.8 (-8.6 to -4.7)*	11.2 (4.2-24.3)	6.7 (4.0 to 9.5)*	-6.6 (-8.9 to -4.1)*
Maternal hypertensive disorders	4624.4 (3030.8 to 6592.4)	2.7 (-37.9 to 69.8)*	-5.8 (-43.0 to 55.4)*	222.4 (118.4-365.4)	2.6 (-37.3 to 68.5)	-5.9 (-42.4 to 54.4)
Maternal hypertensive disorders episodes aggregate	4629.2 (3031.5 to 6602.4)	2.8 (-37.8 to 70.0)	-5.7 (-42.9 to 55.7)	219.1 (115.4-363.3)	2.7 (-37.0 to 68.8)	-5.8 (-42.4 to 55.2)
Eclampsia	5.4 (2.2 to 10.5)	-5.8 (-69.0 to 182.2)	-12.9 (-71.6 to 158.3)	3.3 (1.2-6.3)	-5.7 (-69.0 to 182.2)	-12.8 (-71.6 to 158.3)
Maternal obstructed labour and uterine rupture	1077.2 (933.3 to 1255.2)	-13.1 (-17.5 to -7.9)*	-23.9 (-27.7 to -19.7)*	352.0 (234.6-499.9)	-12.6 (-17.3 to -7.3)*	-23.4 (-27.4 to -18.9)*
Obstructed labour, acute event	89.6 (53.0 to 142.9)	-3.6 (-45.9 to 74.7)	-11.4 (-49.7 to 60.1)	27.7 (14.8-47.0)	-3.1 (-44.9 to 72.1)	-10.9 (-49.0 to 58.2)
Fistula due to maternal obstructed labour and uterine rupture aggregate	987.6 (850.6 to 1156.0)	-13.9 (-16.3 to -11.4)*	-24.9 (-27.0 to -22.8)*	324.3 (216.2-457.4)	-13.3 (-16.6 to -10.1)*	-24.3 (-27.1 to -21.7)*
Maternal abortion, miscarriage, and ectopic pregnancy	441.3 (285.5 to 630.6)	-0.4 (-37.2 to 58.1)	-9.4 (-42.4 to 44.2)	48.2 (27.2-78.5)	-0.4 (-36.9 to 60.5)	-9.3 (-42.4 to 45.6)
Other maternal disorders	135.0 (90.6-190.2)	-2.2 (-19.6 to 17.5)	-12.2 (-27.7 to 5.5)
Neonatal disorders	52 961.9 (50 435.8 to 54 978.4)*	14.7 (11.9 to 17.8)*	3.9 (1.3 to 6.7)*	10 710.5 (8113.9-13 786.1)	13.3 (9.5 to 17.4)*	3.1 (-0.3 to 6.8)
Neonatal preterm birth complications	41 855.4 (36 843.4 to 47 188.1)*	10.6 (6.2 to 15.2)*	-0.4 (-4.3 to 3.8)*	5090.9 (3864.5-6555.6)*	4.4 (-1.1 to 10.1)	-5.2 (-10.1 to 0.2)
Vision loss due to retinopathy of prematurity aggregate	4245.1 (3386.2 to 5265.3)	11.1 (7.2 to 15.1)*	-2.3 (-5.8 to 1.3)	135.5 (83.0-208.8)	13.9 (9.5 to 18.4)	-0.8 (-4.5 to 3.2)
Complications of preterm birth complications aggregate	37 609.0 (33 086.7 to 42 697.0)	10.5 (5.6 to 15.7)*	-0.1 (-4.6 to 4.5)	4955.4 (3756.9-6394.8)	4.1 (-1.4 to 10.0)	-5.3 (-10.4 to 0.2)
Neonatal encephalopathy due to birth asphyxia and trauma	13 681.5 (9 797.3 to 19 759.3)	27.3 (24.6 to 30.0)*	16.2 (13.7 to 18.5)*	3769.4 (2451.7-5809.3)	25.1 (22.0 to 28.3)*	14.2 (11.5 to 17.0)*
Neonatal sepsis and other neonatal infections	139.1 (79.9 to 216.0)	-1.6 (-3.0 to -0.0)*	-6.7 (-8.0 to -5.2)*	17.7 (9.2-31.0)	-1.8 (-3.5 to 0.1)	-6.9 (-8.5 to -5.1)*
Severe infection due to neonatal sepsis and other neonatal infections	130.1 (71.1 to 207.6)	-2.2 (-3.8 to -0.5)*	-7.3 (-8.8 to -5.6)*	16.6 (8.1-30.0)	-2.1 (-3.7 to -0.3)*	-7.1 (-8.7 to -5.4)*

(Table 3 continues on next page)

	Prevalence (thousands)			YLDs (thousands)		
	2015	Percentage change in counts between 2005 and 2015	Percentage change in ASR between 2005 and 2015	2015	Percentage change in counts between 2005 and 2015	Percentage change in ASR between 2005 and 2015
(Continued from previous page)						
Complications of neonatal sepsis and other neonatal infections aggregate	8.9 (7.5 to 10.4)	8.0 (4.3 to 12.7)*	2.4 (-1.1 to 6.9)	1.1 (0.6-1.6)	2.6 (-9.4 to 14.4)	-2.7 (-14.1 to 8.5)
Haemolytic disease and other neonatal jaundice	2042.8 (1825.0 to 2309.0)	20.0 (18.8 to 21.4)*	9.2 (8.0 to 10.3)*	603.0 (451.9-770.8)	17.0 (14.8 to 19.4)*	6.5 (4.4 to 8.6)*
Other neonatal disorders	1229.5 (836.5-1 647.2)	19.9 (0.8 to 41.1)*	8.5 (-8.5 to 27.6)
Nutritional deficiencies	1 482 655.3 (1 477 085.9 to	2.7 (2.2-3.2)*	-7.1 (-7.5-6.7)*	54 632.4 (36 772.8-77 894.3)	-3.5 (-4.9-2.1)*	-11.4 (-12.7-10.3)*
Protein-energy malnutrition	22 834.3 (21 764.0 to 23 985.1)	-4.8 (-10.2 to 0.8)	-12.1 (-17.1 to -6.9)*	2823.3 (1820.3-3984.2)	-4.5 (-9.8 to 1.1)	-11.8 (-16.7 to -6.6)*
Iodine deficiency	110 919.5 (100 337.2 to 125 252.8)	7.0 (4.7 to 9.4)*	-6.0 (-8.0 to -3.8)*	2386.9 (1 508.3-3 564.2)	7.7 (5.7 to 9.8)*	-4.4 (-6.4 to -2.4)*
Visible goiter due to iodine deficiency aggregate	108 307.6 (97 705.7 to 122 260.0)	6.9 (4.5 to 9.4)*	-6.2 (-8.2 to -4.0)*	1927.5 (1192.0-3040.1)	6.9 (4.4 to 9.3)*	-6.0 (-8.2 to -3.9)*
Visible goiter with heart failure due to iodine deficiency aggregate	0.3 (0.3 to 0.4)	39.6 (38.3 to 41.0)*	4.2 (3.3 to 5.2)*	0.0 (0.0-0.1)	39.6 (38.3 to 41.0)*	4.2 (3.3 to 5.2)*
Intellectual disability due to iodine deficiency aggregate	2611.6 (1085.6 to 3621.3)	10.4 (6.1 to 12.6)*	2.0 (-2.0 to 4.3)	459.3 (163.4-752.8)	11.0 (6.4 to 13.6)*	2.8 (-1.6 to 5.2)
Vitamin A deficiency	4901.1 (3975.6 to 5921.7)	9.3 (6.2 to 12.5)*	0.1 (-2.7 to 2.7)	232.4 (143.5-345.9)	10.9 (7.5 to 14.6)*	0.2 (-2.7 to 3.3)
Iron-deficiency anaemia	1 477 530.5 (1 470 902.3 to 1 485 322.2)	1.5 (0.9 to 2.1)*	-8.0 (-8.5 to -7.4)*	48 529.2 (32 560.8-69 725.2)	-3.8 (-5.1 to -2.4)*	-11.6 (-12.8 to -10.5)*
Iron-deficiency anaemia without heart failure aggregate	1 477 458.4 (1 470 824.4 to	39.7 (38.1-41.3)*	6.1 (4.7-7.4)*	8.1 (5.5-11.4)	39.6 (35.2-44.3)*	6.0 (2.5-9.9)*
Iron-deficiency anaemia with heart failure aggregate	72.1 (64.6 to 80.7)	39.7 (38.1 to 41.3)*	6.1 (4.7 to 7.4)*	8.1 (5.5-11.4)	39.6 (35.2 to 44.3)*	6.0 (2.5 to 9.9)*
Other nutritional deficiencies	660.7 (396.0-1026.9)	-16.7 (-36.8 to 5.8)	-22.9 (-41.5 to -2.1)*
Other communicable, maternal, neonatal, and nutritional diseases	1 604 919.9 (1 563 786.8 to 1 636 531.0)	16.1 (15.3 to 17.0)*	-0.4 (-1.1 to 0.4)	3929.7 (2566.9-5786.6)	6.5 (3.9 to 9.2)*	-4.0 (-6.2 to -1.7)*
Sexually transmitted diseases excluding HIV	1 145 527.5 (1 100 971.2 to 1 176 685.1)	16.9 (15.7 to 18.3)*	-1.0 (-2.1 to 0.1)	1573.7 (974.0-2501.8)	16.9 (14.5 to 19.1)*	1.3 (-0.6 to 3.4)
Syphilis	43 604.9 (37 160.3 to 50 658.3)	3.7 (-0.4 to 7.0)	-9.8 (-13.4 to -6.9)*	242.8 (166.1-333.7)	15.3 (12.2 to 18.5)*	-5.6 (-8.0 to -3.0)*
Early syphilis aggregate	42 350.9 (35 898.4 to 49 441.3)	3.3 (-0.9 to 6.8)	-9.9 (-13.6 to -6.9)*	10.3 (3.0-25.3)	3.4 (-1.1 to 7.4)	-9.9 (-13.7 to -6.3)*
Adult tertiary syphilis	1254.0 (1127.1 to 1385.8)	15.8 (13.2 to 18.3)*	-5.6 (-7.8 to -3.6)*	232.6 (155.4-322.8)	15.9 (12.6 to 19.1)*	-5.4 (-7.9 to -2.8)*
Chlamydial infection	82 822.4 (66 426.4 to 104 328.5)	8.3 (6.1 to 10.7)*	-2.9 (-4.8 to -0.9)*	364.6 (210.6-589.5)	10.1 (6.9 to 13.4)*	-1.8 (-4.5 to 0.8)
Chlamydial infection episodes aggregate	81 072.5 (64 735.8 to 102 635.8)	8.3 (6.0 to 10.7)*	-2.9 (-4.8 to -0.9)*	332.7 (185.2-546.0)	10.7 (7.1 to 14.4)*	-1.1 (-4.1 to 2.1)
Pelvic inflammatory diseases due to chlamydial infection aggregate	1.73 (147.8 to 202.0)	1.4 (-0.8 to 3.4)	-11.5 (-13.2 to -9.8)*	23.1 (15.5 to 32.5)	1.8 (-2.2 to 5.7)	-11.1 (-14.5 to 7.8)*

(Table 3 continues on next page)

	Prevalence (thousands)			YLDs (thousands)		
	2015	Percentage change in counts between 2005 and 2015	Percentage change in ASR between 2005 and 2015	2015	Percentage change in counts between 2005 and 2015	Percentage change in ASR between 2005 and 2015
(Continued from previous page)						
Infertility due to chlamydial infection aggregate	1576.6 (1450.8 to 1703.9)	11.1 (10.0 to 12.2)*	-3.0 (-3.9 to -2.0)	8.8 (3.6 to 18.4)	11.1 (9.0 to 13.2)*	-2.9 (-4.8 to -1.1)*
Gonococcal infection	47 468.5 (35 848.1 to 62 099.7)	25.3 (18.3 to 31.2)*	12.2 (5.9 to 17.7)*	444.9 (260.8-691.8)	26.3 (20.0 to 31.4)*	12.6 (6.7 to 17.3)*
Gonococcal infection episodes aggregate	46 561.5 (34 928.2 to 61 227.4)	25.6 (18.5 to 31.6)*	12.5 (6.1 to 18.1)*	430.3 (250.5-671.5)	27.3 (20.7 to 32.5)*	13.5 (7.4 to 18.3)*
Pelvic inflammatory diseases due to gonococcal infection aggregate	907.0 (822.0 to 1 016.3)	10.0 (7.6 to 11.8)*	-3.8 (-6.0 to -2.2)*	14.6 (9.6-21.4)	3.5 (-0.8 to 7.1)	-8.7 (-12.4 to -5.7)*
Trichomoniasis	167 618.6 (144 744.1 to 193 444.0)	16.2 (15.3 to 17.2)*	0.9 (0.4 to 1.3)*	194.3 (78.0-408.3)	16.1 (15.0 to 17.2)*	1.0 (0.3 to 1.8)*
Genital herpes	885 169.4 (772 303.8 to 1 008 588.1)	18.1 (16.4 to 19.9)*	-1.3 (-2.7 to 0.1)	236.4 (74.3-554.2)	19.5 (17.4 to 23.0)*	0.7 (-1.4 to 5.0)
Moderate infection due to initial genital herpes episode	389.0 (88.0 to 909.7)	38.1 (33.6 to 43.2)*	29.5 (25.4 to 34.3)*	19.0 (4.3-45.8)	37.6 (30.9 to 45.6)*	29.1 (22.9 to 36.7)*
Complications of genital herpes aggregate	884 780.3 (771 914.6 to 1 008 400.6)	18.1 (16.4 to 19.9)*	-1.3 (-2.7 to 0.0)	217.3 (62.2-534.5)	18.1 (16.3 to 19.9)*	-1.2 (-2.7 to 0.3)
Other sexually transmitted diseases	4821.8 (4452.8 to 5174.6)	9.6 (8.1 to 11.0)*	-4.7 (-5.9 to -3.6)*	90.7 (61.5-129.7)	3.7 (1.1 to 6.2)*	-10.0 (-12.2 to -7.8)*
Pelvic inflammatory diseases due to other sexually transmitted diseases aggregate	507.2 (437.2 to 591.9)	1.2 (-0.1 to 2.6)	-12.2 (-13.1 to -11.2)*	66.8 (45.4-92.9)	1.3 (-1.2 to 4.2)	-12.0 (-14.3 to -9.8)*
Infertility due to other sexually transmitted diseases aggregate	4314.5 (3943.8 to 4657.1)	10.7 (9.1 to 12.2)*	-3.8 (-5.1 to -2.4)*	23.9 (9.7-48.8)	10.8 (9.0 to 12.7)*	-3.6 (-5.2 to -2.0)*
Other sexually transmitted diseases	329.4 (200.1-504.2)	19.5 (14.4 to 24.6)*	6.7 (2.2 to 11.3)*
Hepatitis	497 901.2 (490 397.2 to 505 815.2)	16.7 (15.8 to 17.6)*	2.2 (1.4 to 2.9)*	406.4 (267.6-588.8)	12.6 (1.7 to 24.1)*	1.2 (-8.3 to 11.5)
Acute hepatitis A	8785.5 (7 950.0 to 9 598.2)	4.2 (-7.2 to 17.0)	-2.8 (-13.5 to 9.1)	172.5 (111.5-249.7)	9.5 (-0.1 to 20.4)	1.9 (-6.9 to 12.0)
Hepatitis B	356 083.4 (342 942.4 to 370 231.5)	16.7 (15.0 to 18.4)*	2.8 (1.3 to 4.3)*	190.2 (121.1-282.2)	16.9 (-4.5 to 40.2)	1.2 (-16.9 to 20.7)
Acute hepatitis B aggregate	12 832.2 (11 239.6 to 14 567.4)	13.2 (-4.3 to 35.2)	1.0 (-14.6 to 20.0)	190.2 (121.1-282.2)	16.9 (-4.5 to 40.2)	1.2 (-16.9 to 20.7)
Chronic hepatitis B	343 251.2 (330 541.3 to 357 194.6)	16.9 (15.3 to 18.4)*	2.8 (1.5 to 4.3)*	0.0 (0.0-0.0)	0.0 (0.0 to 0.0)	0.0 (0.0 to 0.0)
Hepatitis C	142 745.3 (127 559.8 to 157 704.1)	18.0 (16.7 to 19.2)*	1.2 (0.1 to 2.2)*	8.7 (4.3-16.5)	17.0 (12.2 to 21.7)*	3.3 (-0.5 to 7.3)
Acute hepatitis C aggregate	622.3 (580.4 to 666.6)	17.0 (15.7 to 18.4)*	3.4 (2.5 to 4.3)*	8.7 (4.3-16.5)	17.0 (12.2 to 21.7)*	3.3 (-0.5 to 7.3)
Chronic hepatitis C	142 122.9 (126 977.7 to 157 044.9)	18.0 (16.7 to 19.2)*	1.2 (0.1 to 2.2)*	0.0 (0.0-0.0)	0.0 (0.0 to 0.0)	0.0 (0.0 to 0.0)
Acute hepatitis E	1501.9 (1385.4 to 1636.4)	3.5 (2.2 to 4.7)*	-3.2 (-4.1 to -2.3)*	34.9 (22.4-51.7)	4.9 (-4.4 to 15.5)	-2.6 (-10.9 to 6.8)
Other infectious diseases	54 835.4 (54 385.8 to 55 273.8)	0.4 (-1.6 to 2.3)	-8.0 (-9.8 to -6.2)*	1949.6 (1307.1-2797.1)	-1.6 (-5.1 to 2.0)	-8.7 (-12.0 to -5.5)*
Non-communicable diseases	6 652 153.4 (6 633 099.8 to 6 668 805.2)	13.7 (13.6 to 13.8)*	0.0 (-0.0 to 0.1)	638 480.8 (478 716.6-819 498.9)	18.8 (18.4 to 19.2)*	-0.1 (-0.4 to 0.2)

(Table 3 continues on next page)

	Prevalence (thousands)			YLDs (thousands)		
	2015	Percentage change in counts between 2005 and 2015	Percentage change in ASR between 2005 and 2015	2015	Percentage change in counts between 2005 and 2015	Percentage change in ASR between 2005 and 2015
(Continued from previous page)						
Neoplasms	90 497.5 (89 215.7 to 91 896.1)	44.4 (42.1 to 46.7)*	12.4 (10.6 to 14.2)*	8569.3 (6265.5-11 079.2)	36.6 (31.8 to 41.2)*	6.4 (2.7 to 9.7)*
Lip and oral cavity cancer	2425.1 (2278.7 to 2582.3)	38.6 (30.3 to 47.1)*	7.9 (1.5 to 14.3)*	209.1 (150.2-271.4)	36.7 (28.6 to 45.3)*	6.2 (-0.1 to 12.7)
Nasopharynx cancer	732.7 (580.5 to 883.9)	18.0 (0.2 to 39.8)*	-5.5 (-19.2 to 11.1)	69.7 (47.2-95.8)	16.8 (0.8 to 36.0)*	-6.8 (-18.7 to 7.7)
Other pharynx cancer	945.5 (885.9 to 1015.0)	32.7 (23.8 to 41.5)*	2.2 (-4.5 to 9.0)	79.5 (57.8-104.3)	30.7 (21.9 to 39.2)*	0.6 (-6.2 to 7.2)
Oesophageal cancer	746.0 (641.5 to 925.7)	19.6 (2.5 to 39.9)*	-8.2 (-21.2 to 7.1)	128.6 (92.0-167.5)	9.5 (-3.1 to 24.1)	-16.1 (-25.4 to -5.2)*
Stomach cancer	3539.4 (3339.9 to 3776.2)	16.3 (8.9 to 24.4)*	-9.9 (-15.5 to -3.7)*	396.5 (292.4-504.2)	12.0 (4.8 to 20.2)*	-13.3 (-18.8 to -7.1)*
Colon and rectum cancer	9399.0 (9059.3 to 9758.3)	42.6 (38.1 to 47.2)*	8.7 (5.4 to 12.2)*	762.8 (564.2-979.6)	38.4 (33.7 to 43.3)*	5.3 (1.9 to 8.9)*
Colon and rectum cancer aggregate	9377.0 (9037.3 to 9736.4)	42.6 (38.1 to 47.3)*	8.7 (5.4 to 12.2)*	759.8 (561.9-975.3)	38.4 (33.7 to 43.3)*	5.4 (1.9 to 9.0)*
Stoma due to colon and rectum cancer	22.0 (21.3 to 22.8)	26.1 (25.0 to 27.2)*	-4.9 (-5.7 to -4.1)*	3.0 (2.0-4.1)*	26.0 (20.7 to 31.6)*	-4.7 (-9.4 to -0.0)*
Liver cancer	618.7 (550.7 to 688.0)	59.8 (38.2 to 82.2)*	23.6 (6.8 to 40.6)*	188.2 (130.4-245.6)	25.2 (9.9 to 45.0)*	-1.8 (-13.5 to 13.2)
Liver cancer due to hepatitis B	305.9 (235.8 to 418.9)	20.6 (-9.1 to 63.2)	-3.6 (-26.5 to 29.1)	59.9 (40.2-80.4)	9.5 (-8.3 to 34.1)	-12.5 (-26.5 to 6.1)
Liver cancer due to hepatitis C	268.7 (228.4 to 320.8)	80.1 (53.9 to 113.8)*	38.1 (17.1 to 65.1)*	44.3 (31.6-57.3)	47.7 (34.2 to 65.9)*	13.2 (2.7 to 27.7)*
Liver cancer due to alcohol use	273.6 (229.0 to 350.7)	58.8 (28.9 to 94.8)*	22.1 (-0.3 to 49.1)	53.5 (37.3-70.1)	39.3 (22.1 to 61.6)*	7.2 (-5.8 to 23.5)
Liver cancer due to other causes	151.0 (118.8 to 201.0)	24.9 (-5.6 to 63.5)	-2.0 (-26.3 to 29.5)	30.5 (20.7-40.6)	12.1 (-4.5 to 32.8)	-11.9 (-24.9 to 4.8)
Gallbladder and biliary tract cancer	149.4 (138.5 to 158.9)	22.0 (14.4 to 30.1)*	-7.4 (-13.2 to -1.3)*	39.5 (27.7-51.5)	19.0 (11.8 to 26.8)*	-9.5 (-15.1 to -3.7)*
Pancreatic cancer	393.8 (372.9 to 417.1)	46.2 (40.7 to 52.1)*	10.9 (6.7 to 15.4)*	86.6 (61.1-112.2)	38.4 (33.1 to 43.8)*	5.1 (1.1 to 9.3)*
Larynx cancer	1412.6 (1340.0 to 1499.9)	26.2 (20.1 to 33.0)*	-2.3 (-7.0 to 2.9)	147.3 (105.7-194.3)	24.0 (18.1 to 30.0)*	-3.9 (-8.4 to 0.7)
Larynx cancer aggregate	820.9 (749.1 to 911.1)	24.4 (14.2 to 35.8)*	-4.9 (-12.5 to 3.4)	93.6 (67.4-121.5)	21.5 (12.7 to 30.8)*	-6.8 (-13.1 to 0.2)
Laryngectomy due to larynx cancer	591.0 (576.5 to 605.4)	28.7 (28.0 to 29.3)*	1.8 (1.3 to 2.4)*	53.7 (35.8-77.6)	28.6 (25.8 to 31.4)*	1.9 (-0.2 to 4.1)
Tracheal, bronchus, and lung cancer	3299.7 (3095.1 to 3536.4)	37.7 (29.6 to 47.1)*	5.6 (-0.5 to 12.6)	513.6 (377.9-648.8)	31.1 (23.4 to 40.0)*	0.6 (-5.2 to 7.1)
Malignant skin melanoma	3082.3 (2475.8 to 3901.7)	59.0 (50.7 to 67.0)*	26.2 (19.5 to 32.6)*	180.5 (120.8-257.6)	56.3 (47.9 to 64.0)*	23.7 (16.9 to 29.9)*
Non-melanoma skin cancer	2558.2 (2494.8 to 2625.6)	76.8 (72.0 to 81.6)*	34.5 (30.8 to 38.2)*	148.0 (104.7-197.7)	82.9 (72.7 to 93.9)*	40.1 (31.9 to 48.7)*
Non-melanoma skin cancer (squamous-cell carcinoma)	2155.9 (2019.5 to 2312.2)	93.5 (82.2 to 105.8)*	49.6 (40.5 to 59.2)*	142.0 (100.7-187.1)	86.4 (75.6 to 98.2)*	42.8 (34.2 to 52.1)*
Non-melanoma skin cancer (basal-cell carcinoma)	578.1 (497.7 to 671.7)	27.0 (24.0 to 29.7)*	-3.9 (-6.0 to -1.8)*	6.0 (2.9-11.1)	26.9 (23.5 to 30.1)*	-3.8 (-6.4 to -1.2)*
Breast cancer	21361.8 (20 249.5 to 22 266.3)	41.5 (33.4 to 49.6)*	9.9 (4.1 to 15.7)*	1796.5 (1270.7-2411.6)	36.1 (29.2 to 42.7)*	5.3 (0.4 to 10.1)*
Breast cancer aggregate	10718.5 (9581.7 to 11 667.2)	62.7 (42.9 to 83.3)*	32.0 (16.2 to 48.1)*	989.8 (719.8-1271.0)	46.6 (32.7 to 61.5)*	15.8 (5.3 to 27.5)*
Mastectomy due to breast cancer	10638.5 (10 404.8 to 10 894.5)	25.1 (24.1 to 26.1)*	-4.7 (-5.4 to -3.9)*	806.8 (520.3-1171.8)	25.1 (24.0 to 26.2)*	-4.6 (-5.5 to -3.8)*

(Table 3 continues on next page)

	Prevalence (thousands)			YLDs (thousands)		
	2015	Percentage change in counts between 2005 and 2015	Percentage change in ASR between 2005 and 2015	2015	Percentage change in counts between 2005 and 2015	Percentage change in ASR between 2005 and 2015
(Continued from previous page)						
Cervical cancer	3442.4 (3129.9 to 3765.0)	-1.6 (-11.4 to 9.8)	-20.5 (-28.2 to -11.5)*	264.0 (187.9-342.6)	-1.4 (-10.9 to 9.8)	-20.5 (-28.0 to -11.6)*
Uterine cancer	3827.7 (3425.6 to 4285.2)	39.5 (22.9 to 55.9)*	8.2 (-4.0 to 20.0)	250.2 (172.4-341.0)	37.1 (21.1 to 52.9)*	6.1 (-5.4 to 17.6)
Ovarian cancer	1174.2 (1107.4 to 1250.1)	28.3 (20.8 to 36.3)*	0.3 (-5.1 to 6.3)	150.7 (110.5-192.3)	25.8 (17.8 to 33.9)*	-1.7 (-7.4 to 4.3)
Prostate cancer	14 434.4 (11 932.2 to 19 785.1)	70.6 (61.6 to 81.4)*	29.7 (22.5 to 38.4)*	1150.3 (804.9-1 643.3)	60.5 (51.7 to 70.6)*	21.5 (14.5 to 29.6)*
Prostate cancer aggregate	13 492.3 (10 984.7 to 18 837.5)	74.6 (64.7 to 86.3)*	32.5 (24.6 to 42.1)*	1069.7 (739.5-1 543.7)	63.5 (53.7 to 74.9)*	23.5 (15.8 to 32.6)*
Impotence and incontinence due to prostate cancer aggregate	942.0 (927.1 to 957.3)	29.0 (28.6 to 29.4)*	-0.9 (-1.2 to -0.5)*	80.6 (55.2-111.8)	28.9 (27.1 to 30.5)*	-1.0 (-2.4 to 0.2)
Testicular cancer	685.8 (634.7 to 732.4)	41.9 (30.8 to 52.4)*	23.4 (13.9 to 32.2)*	42.1 (28.4-58.3)	40.3 (29.3 to 50.6)*	21.5 (11.9 to 30.4)*
Kidney cancer	2870.3 (2728.0 to 3031.5)	57.9 (50.0 to 66.0)*	22.8 (16.8 to 29.1)*	202.7 (145.7-270.5)	54.4 (46.9 to 61.8)*	19.8 (14.0 to 25.6)*
Bladder cancer	3407.9 (3240.0 to 3603.3)	35.7 (28.6 to 43.3)*	3.3 (-2.0 to 9.0)	267.0 (193.7-349.4)	32.1 (25.8 to 38.8)*	0.7 (-4.0 to 5.6)
Bladder cancer episodes aggregate	3273.4 (3 105.3 to 3 470.7)	36.3 (28.9 to 44.3)*	3.7 (-1.8 to 9.6)	244.1 (176.2-317.4)	33.1 (26.1 to 40.4)*	1.2 (-3.9 to 6.7)
Urinary incontinence due to bladder cancer	134.3 (129.7 to 139.4)	22.8 (22.0 to 23.7)*	-5.7 (-6.2 to -5.1)*	22.9 (15.9-31.2)	23.0 (18.7 to 27.1)*	-5.3 (-8.6 to -2.2)*
Brain and nervous system cancer	1205.1 (1101.7 to 1322.9)	25.3 (14.9 to 37.6)*	4.2 (-3.9 to 13.8)	126.1 (90.3-164.5)	24.7 (15.0 to 36.3)*	2.8 (-4.7 to 12.0)
Thyroid cancer	3166.5 (2936.7 to 3340.6)	101.1 (86.4 to 112.9)*	60.8 (48.9 to 70.2)*	190.0 (132.5-259.3)	96.8 (81.6 to 109.2)*	56.5 (44.3 to 66.3)*
Mesothelioma	60.8 (58.1 to 63.6)	41.3 (35.1 to 47.7)*	9.5 (4.6 to 14.5)*	12.2 (8.7-15.8)	39.5 (32.1 to 46.8)*	7.9 (2.1 to 13.9)*
Hodgkin's lymphoma	574.4 (519.0 to 672.9)	17.8 (10.5 to 25.3)*	0.0 (-6.1 to 6.4)	48.5 (33.7-65.8)	14.6 (7.4 to 21.7)*	-3.6 (-9.6 to 2.4)*
Non-Hodgkin lymphoma	4292.3 (3741.0 to 4593.6)	63.5 (47.5 to 74.3)*	30.5 (19.1 to 38.4)*	312.3 (221.8-414.2)	57.9 (43.8 to 67.8)*	25.6 (15.1 to 33.1)*
Multiple myeloma	488.2 (449.4 to 527.7)	54.6 (44.8 to 64.5)*	18.0 (10.7 to 25.4)*	104.0 (73.6-134.8)	49.0 (40.0 to 58.5)*	13.6 (6.9 to 21.0)*
Leukaemia	2303.6 (2216.1 to 2384.5)	35.8 (29.7 to 41.8)*	10.2 (5.7 to 14.8)*	379.4 (276.2-488.9)	28.6 (21.2 to 36.9)*	4.3 (-1.1 to 10.4)
Acute lymphoid leukaemia	875.5 (709.0 to 1072.3)	25.4 (7.3 to 46.4)*	10.2 (-5.0 to 27.8)	97.3 (68.0-129.8)	25.7 (11.5 to 40.8)*	8.1 (-3.1 to 20.0)
Chronic lymphoid leukaemia	904.0 (833.2 to 974.4)	31.7 (22.2 to 42.9)*	4.5 (-2.2 to 12.4)	119.8 (87.2-152.9)	27.6 (18.9 to 37.4)*	0.5 (-6.0 to 7.7)
Acute myeloid leukaemia	999.3 (882.5 to 1136.8)	38.4 (26.6 to 50.7)*	14.2 (5.7 to 23.5)*	120.9 (87.3-155.6)	36.5 (27.5 to 45.6)*	10.4 (3.7 to 17.4)*
Chronic myeloid leukaemia	297.7 (271.8 to 324.7)	22.2 (14.3 to 31.0)*	-3.4 (-9.3 to 3.4)	41.4 (29.5-53.6)	17.7 (10.2 to 26.0)*	-7.5 (-13.3 to -1.2)*
Other neoplasms	4577.5 (4145.5 to 4922.0)	36.1 (27.2 to 44.5)*	10.9 (4.1 to 17.6)*	323.2 (228.6-431.8)	34.4 (26.2 to 42.6)*	9.0 (2.4 to 15.4)*
Cardiovascular diseases	422 738.4 (415 534.5 to 427 870.8)	24.8 (24.0 to 25.6)*	-1.8 (-2.5 to -1.2)*	25 620.1 (18 401.6-33 656.6)	23.5 (21.7 to 24.5)*	-2.2 (-3.2 to -1.7)*
Rheumatic heart disease	33 438.8 (29 725.7 to 43 119.8)	-4.7 (-11.0 to 0.1)	-16.3 (-22.5 to -12.1)*	1654.7 (1041.4-2530.9)	-3.4 (-9.9 to 1.4)	-15.4 (-21.5 to -11.1)*
Rheumatic heart disease, without heart failure	32 236.8 (28 520.8 to 41 912.7)	-5.6 (-11.9 to -0.8)*	-16.9 (-23.1 to -12.7)*	1518.6 (941.1-2356.4)	-5.5 (-11.7 to -0.6)*	-16.8 (-22.9 to -12.3)*

(Table 3 continues on next page)

(Continued from previous page)

	Prevalence (thousands)			YLDs (thousands)		
	2015	Percentage change in counts between 2005 and 2015	Percentage change in ASR between 2005 and 2015	2015	Percentage change in counts between 2005 and 2015	Percentage change in ASR between 2005 and 2015
Heart failure due to rheumatic heart disease aggregate	1202.0 (1138.2 to 1269.0)	29.8 (28.5 to 31.2)*	2.2 (1.3 to 3.2)*	136.1 (94.5-187.6)	29.6 (27.2 to 32.1)*	2.4 (0.6 to 4.2)*
Ischaemic heart disease	110 550.3 (100 675.9 to 121 798.7)	25.9 (24.6 to 27.2)*	-3.4 (-4.2 to -2.6)*	7274.7 (4958.6-9940.3)	30.2 (29.1 to 31.3)*	-0.3 (-1.0 to 0.3)
Myocardial infarction episodes aggregate	15 930.1 (14 106.5 to 17 764.0)	6.4 (1.9 to 10.6)*	-18.1 (-21.6 to -14.8)*	33.0 (23.0-44.9)	15.2 (12.6 to 17.9)*	-13.1 (-15.1 to -11.1)*
Angina due to ischaemic heart disease aggregate	72 344.6 (62 699.8 to 83 460.0)	29.3 (27.7 to 30.7)*	-0.3 (-1.2 to 0.5)	4794.5 (3100.7-6633.6)	29.5 (27.8 to 31.0)*	-0.0 (-1.0 to 0.8)
Heart failure due to ischaemic heart disease aggregate	22 275.7 (21 272.0 to 23 320.4)	31.9 (31.1 to 32.7)*	-0.6 (-1.2 to -0.1)*	2447.2 (1736.8-3342.6)	31.9 (31.0 to 32.8)*	-0.5 (-1.1 to 0.1)
Cerebrovascular disease	42 430.9 (42 068.2 to 42 767.1)	21.0 (20.4 to 21.5)*	-4.4 (-4.8 to -3.9)*	6455.2 (4487.0-8609.6)	20.7 (19.8 to 21.6)*	-4.2 (-4.9 to -3.5)*
Ischaemic stroke	24 929.0 (24 362.2 to 25 610.0)	21.8 (21.0 to 22.6)*	-5.2 (-5.8 to -4.6)*	3660.0 (2559.3-4874.9)	22.0 (21.0 to 23.0)*	-4.9 (-5.7 to -4.2)*
Chronic ischaemic stroke aggregate	24 663.2 (24 092.1 to 25 345.8)	21.9 (21.1 to 22.7)*	-5.1 (-5.8 to -4.5)*	3605.1 (2512.6-4815.3)	22.1 (21.1 to 23.1)*	-4.9 (-5.6 to -4.1)*
Ischaemic stroke episodes aggregate	265.8 (246.6 to 285.1)	16.8 (14.2 to 19.7)*	-10.6 (-12.7 to -8.5)*	54.9 (36.3-74.5)	17.5 (13.8 to 20.9)*	-10.0 (-12.9 to -7.2)*
Haemorrhagic stroke	18 669.6 (18 258.7 to 19 124.5)	19.2 (18.5 to 19.9)*	-3.3 (-3.8 to -2.7)*	2795.2 (1945.8-3749.6)	19.1 (18.1 to 20.2)*	-3.1 (-3.9 to -2.3)*
Chronic haemorrhagic stroke aggregate	18 494.8 (18 088.5 to 18 959.5)	19.3 (18.6 to 19.9)*	-3.2 (-3.8 to -2.6)*	2757.3 (1918.2-3700.6)	19.2 (18.1 to 20.3)*	-3.0 (-3.8 to -2.2)*
Acute haemorrhagic stroke aggregate	174.8 (162.3 to 187.8)	14.3 (11.6 to 16.6)*	-9.1 (-11.1 to -7.3)*	37.9 (24.7-51.2)	14.6 (11.8 to 17.0)*	-8.8 (-11.0 to -6.9)*
Hypertensive heart disease	6086.2 (5732.7 to 6434.1)	37.1 (36.2 to 37.9)*	3.4 (2.9 to 3.9)*	670.0 (467.1-917.1)	37.2 (35.9 to 38.4)*	3.6 (2.8 to 4.4)*
Cardiomyopathy and myocarditis	2536.8 (2404.8 to 2661.4)	26.7 (25.6 to 27.7)*	-1.3 (-2.0 to -0.7)*	275.1 (190.3-377.9)	26.3 (24.6 to 28.1)*	-1.3 (-2.5 to -0.2)*
Acute myocarditis	156.3 (137.1 to 179.6)	21.4 (19.1 to 23.7)*	2.2 (1.3 to 3.1)*	7.8 (4.8-11.6)	21.1 (18.5 to 23.9)*	2.1 (0.3 to 3.8)*
Heart failure due to cardiomyopathy aggregate	2380.4 (2254.3 to 2502.2)	27.0 (25.9 to 28.1)*	-1.5 (-2.2 to -0.9)*	267.3 (185.9-367.9)	26.5 (24.7 to 28.3)*	-1.4 (-2.6 to -0.2)*
Atrial fibrillation and flutter	33 294.3 (29 959.8 to 37 202.0)	28.2 (27.2 to 29.1)*	-2.5 (-3.1 to -1.9)*	2634.6 (1782.6-3637.3)	28.2 (27.1 to 29.3)*	-2.4 (-3.1 to -1.7)*
Peripheral vascular disease	154 650.6 (136 318.0 to 176 210.9)	34.4 (33.4 to 35.2)*	1.8 (1.3 to 2.3)*	572.8 (272.1-1056.2)	34.5 (32.4 to 36.8)*	2.0 (1.2 to 2.8)*
Endocarditis	115.7 (108.1 to 124.8)	22.7 (20.9 to 24.5)*	-5.4 (-6.7 to -4.1)*	12.3 (8.5-17.1)	22.9 (18.6 to 27.6)*	-5.2 (-8.8 to -1.2)*
Endocarditis episodes aggregate	16.9 (15.7 to 18.2)	22.8 (20.3 to 25.3)*	-0.7 (-2.7 to 1.2)	1.0 (0.6-1.4)	22.7 (20.0 to 25.3)*	-0.9 (-3.1 to 1.2)
Heart failure due to endocarditis aggregate	98.8 (91.1 to 107.6)	22.7 (20.7 to 24.7)*	-6.1 (-7.6 to -4.6)*	11.4 (7.9-15.9)	22.9 (18.3 to 28.0)*	-5.5 (-9.3 to -1.2)*
Heart failure due to other cardiovascular diseases aggregate	1126.5 (1045.9 to 1206.1)	34.4 (33.3 to 35.6)*	1.2 (0.4 to 2.0)*	123.7 (86.0-173.3)	34.4 (32.5 to 36.5)*	1.3 (-0.2 to 2.7)
Other cardiovascular diseases episodes aggregate	89 221.1 (83 635.8 to 95 470.4)	24.0 (22.6 to 25.4)*	0.4 (-0.2 to 1.0)*	5947.0 (4074.6-8105.9)	23.7 (22.2 to 25.2)*	0.5 (-0.2 to 1.1)
Chronic respiratory diseases	514 625.8 (503 322.3 to 527 617.0)	12.1 (11.1 to 13.1)*	-3.3 (-4.1 to -2.4)*	30 465.9 (23 341.4-38 294.2)	11.8 (10.2 to 13.6)*	-4.6 (-6.2 to -3.1)*
Chronic obstructive pulmonary disease	174 483.1 (160 204.9 to 188 951.7)	17.0 (15.1 to 19.0)*	-5.8 (-7.3 to -4.4)*	12 047.0 (10 206.8-13 725.4)	16.2 (13.4 to 18.8)*	-5.9 (-8.0 to -3.9)*

(Table 3 continues on next page)

	Prevalence (thousands)			YLDs (thousands)		
	2015	Percentage change in counts between 2005 and 2015	Percentage change in ASR between 2005 and 2015	2015	Percentage change in counts between 2005 and 2015	Percentage change in ASR between 2005 and 2015
(Continued from previous page)						
Chronic obstructive pulmonary disease without heart failure aggregate	168 694.6 (154 443.1 to 182 957.1)	16.5 (14.5 to 18.5)*	-6.1 (-7.6 to -4.6)*	9671.6 (8183.6-11 080.9)	12.3 (9.2 to 15.3)*	-8.0 (-10.5 to -5.6)*
Severe chronic obstructive pulmonary disease with heart failure aggregate	5876.4 (5073.7 to 6548.3)	34.9 (31.1 to 37.8)*	2.4 (-0.3 to 4.2)	2375.5 (1890.3-2747.2)	35.1 (31.4 to 38.2)*	2.5 (-0.0 to 4.5)
Pneumoconiosis	2405.8 (2317.2 to 2486.3)	30.3 (27.8 to 32.9)*	8.2 (6.2 to 10.2)*	474.0 (305.6-682.6)	26.6 (23.6 to 29.3)*	6.0 (3.6 to 8.4)*
Silicosis	402.9 (364.9 to 442.2)	18.1 (15.7 to 20.3)*	-4.3 (-6.1 to -2.6)*	65.7 (41.7-95.4)	18.5 (14.5 to 22.4)*	-3.9 (-7.0 to -0.9)*
Silicosis without heart failure aggregate	389.0 (351.0 to 426.8)	17.6 (15.2 to 19.9)*	-4.6 (-6.4 to -2.8)*	59.2 (36.8-87.9)	17.1 (12.7 to 21.3)*	-4.7 (-8.0 to -1.5)*
Severe silicosis with heart failure aggregate	13.9 (12.0 to 15.6)	32.2 (29.1 to 34.8)*	1.5 (-0.5 to 3.3)	6.4 (4.5-8.5)	33.1 (26.9 to 40.5)*	2.3 (-2.7 to 7.9)
Asbestosis	157.3 (144.8 to 170.9)	30.6 (28.8 to 32.6)*	2.7 (1.3 to 4.1)*	24.8 (16.0-35.6)	30.2 (27.0 to 33.6)*	2.5 (-0.1 to 5.2)
Asbestosis without heart failure aggregate	154.5 (141.8 to 168.1)	30.6 (28.7 to 32.5)*	2.7 (1.3 to 4.1)*	23.5 (15.0-34.0)	29.9 (26.4 to 33.4)*	2.5 (-0.2 to 5.4)
Severe asbestosis with heart failure aggregate	2.8 (2.6 to 3.0)	35.6 (34.2 to 37.1)*	1.1 (0.1 to 2.1)*	1.3 (0.9-1.7)*	35.7 (31.3 to 40.0)*	1.3 (-2.2 to 4.7)
Coal workers pneumoconiosis	84.2 (76.7 to 93.6)	37.0 (34.1 to 39.9)*	7.6 (5.2 to 10.0)*	13.3 (8.4-19.1)*	36.3 (29.4 to 43.4)*	7.0 (1.3 to 12.9)*
Coal workers pneumoconiosis without heart failure aggregate	82.3 (74.9 to 91.7)	36.9 (34.0 to 39.8)*	7.5 (5.2 to 10.0)*	12.4 (7.8-17.9)*	36.0 (28.6 to 43.5)*	6.9 (0.9 to 13.2)*
Severe coal workers pneumoconiosis with heart failure aggregate	1.9 (1.6 to 2.1)	41.6 (37.8 to 45.1)*	8.2 (5.4 to 10.9)*	0.9 (0.6-1.2)*	41.6 (37.7 to 45.2)*	8.2 (5.2 to 10.9)*
Other pneumoconiosis	2388.6 (2164.7 to 2629.1)	27.9 (25.3 to 30.7)*	8.3 (6.0 to 10.5)*	370.2 (239.2-533.9)*	27.5 (23.9 to 31.0)*	8.3 (5.3 to 11.2)*
Other pneumoconiosis without heart failure aggregate	2374.2 (2150.3 to 2615.1)	27.8 (25.2 to 30.7)*	8.3 (6.0 to 10.6)*	363.6 (234.0-524.9)*	27.3 (23.6 to 30.8)*	8.4 (5.3 to 11.4)*
Severe other pneumoconiosis with heart failure aggregate	14.4 (13.1 to 15.6)	41.1 (39.0 to 43.0)*	4.5 (3.4 to 5.5)*	6.5 (4.6-8.5)	40.5 (35.1 to 45.7)*	4.3 (0.3 to 8.4)*
Asthma	358 197.9 (323 133.7 to 393 465.6)	9.5 (7.6 to 11.6)*	-2.5 (-4.3 to -0.5)*	15 898.9 (10 371.0-22 344.1)	9.4 (7.4 to 11.5)*	-2.3 (-4.3 to -0.3)*
Asymptomatic asthma	108 461.1 (92 842.5 to 126 250.5)	9.5 (7.6 to 11.6)*	-2.5 (-4.3 to -0.5)*	0.0 (0.0-0.0)	0.0 (0.0 to 0.0)	0.0 (0.0 to 0.0)
Symptomatic asthma aggregate	249 736.8 (220 469.6 to 278 419.7)	9.5 (7.6 to 11.6)*	-2.5 (-4.3 to -0.5)*	15 898.9 (10 371.0-22 344.1)	9.4 (7.4 to 11.5)*	-2.3 (-4.3 to -0.3)*
Interstitial lung disease and pulmonary sarcoidosis	1916.0 (1757.4 to 2075.5)	25.8 (23.8 to 27.7)*	0.6 (-0.9 to 2.1)	234.7 (147.5-334.2)	25.7 (23.5 to 28.1)*	0.7 (-1.1 to 2.6)
Interstitial lung disease and pulmonary sarcoidosis without heart failure aggregate	1718.7 (1565.7 to 1881.0)	25.2 (23.1 to 27.2)*	0.5 (-1.1 to 2.1)	148.5 (88.8-221.4)	22.7 (19.6 to 25.9)*	0.2 (-2.2 to 2.6)

(Table 3 continues on next page)

	Prevalence (thousands)			YLDs (thousands)		
	2015	Percentage change in counts between 2005 and 2015	Percentage change in ASR between 2005 and 2015	2015	Percentage change in counts between 2005 and 2015	Percentage change in ASR between 2005 and 2015
(Continued from previous page)						
Severe interstitial lung disease and pulmonary sarcoidosis with heart failure aggregate	197.4 (144.1 to 242.7)	31.5 (29.7 to 33.2)*	1.3 (-0.1 to 2.9)	86.2 (54.8-117.6)	31.3 (28.6 to 34.1)*	1.3 (-1.0 to 3.7)
Other chronic respiratory diseases	1811.3 (1483.7-2123.4)	1.7 (-7.3 to 12.2)	-15.8 (-23.5 to -6.9)*
Cirrhosis and other chronic liver diseases	2828.3 (2791.3 to 2862.3)*	34.0 (32.4 to 35.8)*	8.2 (7.0 to 9.5)*	501.0 (352.9-683.1)	31.3 (29.8 to 32.9)*	7.3 (6.1 to 8.4)*
Cirrhosis and other chronic liver diseases due to hepatitis B	796.5 (737.6 to 857.3)	30.5 (28.4 to 32.7)*	4.7 (3.2 to 6.2)*	130.6 (88.7-178.7)	30.5 (27.0 to 33.8)*	4.8 (2.2 to 7.3)*
Cirrhosis and other chronic liver diseases due to hepatitis C	680.8 (628.3 to 734.2)	35.3 (33.0 to 37.5)*	7.6 (6.0 to 9.1)*	111.4 (78.2-152.6)	35.2 (31.6 to 38.6)*	7.6 (5.0 to 10.3)*
Cirrhosis and other chronic liver diseases due to alcohol use	815.6 (758.7 to 877.2)	37.3 (35.4 to 39.3)*	9.8 (8.5 to 11.2)*	133.9 (92.5-185.1)	37.2 (33.9 to 40.5)*	9.9 (7.5 to 12.4)*
Cirrhosis and other chronic liver diseases due to other causes	750.3 (714.3 to 785.9)	23.8 (22.5 to 25.1)*	6.8 (5.9 to 7.6)*	125.0 (87.5-172.2)	23.3 (20.1 to 26.8)*	6.8 (4.2 to 9.5)*
Digestive diseases	258 248.2 (255 386.4 to 260 843.9)	12.3 (10.9 to 13.7)*	-9.4 (-10.4 to -8.3)*	12 142.8 (8 492.0-16 592.5)	11.4 (9.8 to 13.2)*	-9.3 (-10.6 to -7.9)*
Peptic ulcer disease	72 044.2 (71 302.7 to 72 763.7)	4.8 (3.7 to 6.0)*	-17.0 (-17.8 to -16.0)*	2341.7 (1605.6-3293.6)	1.1 (-0.2 to 2.5)	-19.5 (-20.6 to -18.4)*
Peptic ulcer disease symptomatic episodes	4977.7 (4577.6 to 5379.5)	4.7 (2.4 to 6.9)*	-16.7 (-18.4 to -15.0)*	518.7 (357.9-703.2)	4.5 (2.2 to 6.8)*	-16.7 (-18.5 to -14.9)*
Anaemia due to peptic ulcer disease aggregate	67 066.6 (66 465.4 to 67 616.9)	4.8 (3.7 to 6.1)*	-17.0 (-17.9 to -16.0)*	1823.1 (1225.8-2621.2)	0.2 (-1.4 to 1.9)	-20.3 (-21.6 to -19.0)*
Gastritis and duodenitis	161 001.1 (157 928.3 to 164 071.1)	15.5 (13.0 to 17.9)*	-6.9 (-8.8 to -5.0)*	4913.0 (3347.3-6967.1)	11.2 (8.0 to 14.6)*	-9.5 (-11.9 to -6.9)*
Gastritis and duodenitis, symptomatic episodes	3941.0 (3547.0 to 4358.9)	15.4 (10.9 to 17.3)*	-6.9 (-10.8 to -5.7)*	416.0 (277.1-567.9)	15.4 (11.2 to 17.5)*	-6.7 (-10.6 to -5.2)*
Anaemia due to gastritis and duodenitis aggregate	157 060.1 (154 055.1 to 160 141.0)	15.5 (12.9 to 17.9)*	-6.9 (-8.9 to -5.0)*	4497.1 (3050.6-6393.0)	10.8 (7.4 to 14.5)*	-9.8 (-12.4 to -7.0)*
Appendicitis	442.1 (414.6 to 472.3)	14.3 (13.4 to 15.2)*	2.8 (2.1 to 3.4)*	136.0 (92.0-184.6)	14.4 (10.6 to 18.0)*	3.0 (-0.2 to 6.1)
Paralytic ileus and intestinal obstruction	119.7 (110.7 to 130.4)	18.9 (16.9 to 20.7)*	-1.3 (-2.8 to -0.0)*	37.3 (25.1-51.1)	18.6 (15.9 to 21.2)*	-1.1 (-3.4 to 1.2)
Inguinal, femoral, and abdominal hernia	18 476.7 (16 577.2 to 20 339.8)	6.5 (4.2 to 8.6)*	-10.6 (-12.4 to -9.0)*	195.8 (97.0-363.1)	6.5 (4.3 to 8.7)*	-10.4 (-12.2 to -8.8)*
Inflammatory bowel disease	11 223.5 (10 396.8 to 12 000.4)	15.7 (15.0 to 16.3)*	-3.7 (-4.1 to -3.4)*	2387.1 (1653.2-3 263.5)	15.6 (14.7 to 16.4)*	-3.6 (-4.2 to -3.0)*
Vascular intestinal disorders	35.1 (32.4 to 38.1)	23.3 (21.0 to 25.8)*	-2.4 (-4.1 to -0.6)*	11.0 (7.4-14.8)	23.4 (19.7 to 27.2)*	-2.0 (-5.2 to 1.2)
Gallbladder and biliary diseases	5656.5 (5199.3 to 6168.1)	16.9 (14.5 to 19.2)*	-4.1 (-5.9 to -2.2)*	601.0 (409.4-826.4)	16.8 (14.2 to 19.3)*	-3.9 (-5.9 to -1.9)*
Pancreatitis	1018.2 (941.6 to 1 097.4)	24.2 (21.7 to 26.5)*	1.9 (-0.0 to 3.7)	300.0 (203.4-406.7)	24.0 (20.3 to 27.4)*	2.0 (-0.8 to 4.6)
Other digestive diseases	1219.9 (838.5-1677.5)	21.9 (14.4 to 32.4)*	-1.0 (-7.1 to 7.4)
Neurological disorders	2 261 316.4 (2 204 366.6 to 2 316 282.6)	15.5 (14.6 to 16.4)*	0.5 (-0.3 to 1.2)	59 325.6 (40 845.0-81 083.3)	15.6 (14.3 to 16.9)*	-1.4 (-2.4 to -0.4)*
Alzheimer's disease and other dementias	45 956.3 (40 178.5 to 52 655.9)	37.7 (36.2 to 39.0)*	1.1 (0.3 to 1.8)*	6851.2 (4883.9-9062.8)	38.8 (37.1 to 40.5)*	1.1 (0.2 to 1.8)*

(Table 3 continues on next page)

	Prevalence (thousands)			YLDs (thousands)		
	2015	Percentage change in counts between 2005 and 2015	Percentage change in ASR between 2005 and 2015	2015	Percentage change in counts between 2005 and 2015	Percentage change in ASR between 2005 and 2015
(Continued from previous page)						
Parkinson's disease	6193.3 (5725.7 to 6777.2)	31.6 (28.2 to 35.5)*	0.6 (-1.9 to 3.5)	737.8 (516.1-990.1)	31.4 (27.8 to 35.3)*	0.7 (-2.1 to 3.7)
Epilepsy	23 414.5 (21 549.6 to 25 419.4)	11.3 (7.0 to 15.5)*	-1.1 (-5.0 to 2.7)	6286.9 (4315.8-8250.2)	-6.4 (-11.5 to -1.2)*	-16.3 (-20.9 to -11.6)*
Multiple sclerosis	2012.0 (1865.7 to 2166.9)	19.1 (17.0 to 21.3)*	-2.0 (-3.9 to -0.3)*	667.5 (472.0-855.6)	18.8 (16.2 to 21.5)*	-2.0 (-4.2 to 0.1)
Motor neuron disease	202.4 (189.7 to 216.0)	22.3 (19.9 to 24.8)*	-1.3 (-3.3 to 0.7)	42.3 (30.5-55.1)	22.1 (19.5 to 24.8)*	-1.4 (-3.5 to 0.8)
Migraine	958 789.2 (872 109.0 to 1 055 630.6)	15.3 (14.0 to 16.6)*	0.6 (-0.3 to 1.5)	32 898.8 (20 303.6-48 883.2)	15.3 (14.0 to 16.6)*	0.8 (-0.1 to 1.8)
Tension-type headache	1 505 892.3 (1 337 310.0 to 1 681 575.0)	15.3 (14.0 to 16.6)*	0.5 (-0.1 to 1.1)	2260.5 (1058.1-4169.7)	15.3 (14.0 to 16.7)*	0.6 (-0.1 to 1.3)
Medication overuse headache	58 454.5 (50 834.9 to 67 363.9)	19.0 (15.4 to 22.7)*	0.4 (-2.4 to 3.3)	9164.7 (6094.5-13 078.0)	18.9 (15.4 to 22.8)*	0.6 (-2.3 to 3.5)
Other neurological disorders	11.5 (9.1 to 14.6)	17.0 (14.8 to 19.6)*	-0.2 (-0.9 to 0.4)	415.9 (301.2-547.5)	32.2 (27.5 to 36.3)*	-2.1 (-5.3 to 0.9)
Mental and substance use disorders	1 058 903.8 (1 038 544.9 to 1 080 413.8)	14.3 (13.6 to 14.9)*	0.3 (-0.3 to 0.9)	149 977.9 (108 716.1-193 130.8)	16.1 (15.5 to 16.8)*	1.1 (0.7 to 1.4)*
Schizophrenia	23 383.0 (20 608.0 to 26 418.4)	19.5 (18.6 to 20.4)*	0.1 (-0.4 to 0.6)	15 020.5 (10 816.1-18 623.2)	19.5 (18.5 to 20.4)*	0.3 (-0.4 to 0.9)
Alcohol use disorders	63 469.5 (57 507.8 to 69 863.5)	11.1 (8.9 to 13.5)*	-4.6 (-6.5 to -2.4)*	6321.3 (4205.6-8985.8)*	11.1 (9.0 to 13.6)*	-4.5 (-6.4 to -2.3)*
Alcohol dependence aggregate	62 575.3 (56 621.7 to 68 944.3)	11.1 (8.9 to 13.5)*	-4.6 (-6.6 to -2.5)*	6276.2 (4167.7-8929.7)	11.1 (8.9 to 13.6)*	-4.5 (-6.5 to -2.3)*
Fetal alcohol syndrome aggregate	894.2 (833.3 to 951.9)	11.1 (10.3 to 11.7)*	-0.5 (-1.1 to 0.2)	45.1 (29.2-65.1)	10.7 (8.2 to 13.1)*	-0.6 (-2.7 to 1.6)
Drug use disorders	46 388.6 (45 252.4 to 47 446.6)	16.3 (15.0 to 17.6)*	3.9 (2.8 to 5.0)*	9849.4 (6959.0-12775.2)	23.6 (21.5 to 25.9)*	8.2 (6.2 to 10.2)*
Opioid use disorders	16 746.5 (14 659.3 to 19 107.5)	23.3 (20.7 to 26.0)*	6.4 (4.2 to 8.6)*	6969.6 (4842.5-8999.8)	23.3 (20.6 to 26.2)*	6.5 (4.2 to 9.0)*
Asymptomatic opioid dependence	2716.8 (1979.9 to 3704.4)	23.3 (20.7 to 26.0)*	6.4 (4.2 to 8.6)*	0.0 (0.0-0.0)	0.0 (0.0 to 0.0)	0.0 (0.0 to 0.0)*
Symptomatic opioid dependence aggregate	14 029.7 (12 042.2 to 16 122.8)	23.3 (20.7 to 26.0)*	6.4 (4.2 to 8.6)*	6969.6 (4842.5-8999.8)	23.3 (20.6 to 26.2)*	6.5 (4.2 to 9.0)
Cocaine use disorders	3846.3 (3401.6 to 4309.8)	31.1 (28.0 to 34.3)*	14.1 (12.0 to 16.4)*	521.5 (329.0-733.9)	30.6 (27.2 to 34.4)*	14.0 (11.4 to 17.0)*
Asymptomatic cocaine dependence	1927.8 (1544.3 to 2357.8)	31.1 (28.0 to 34.3)*	14.1 (12.0 to 16.4)*	0.0 (0.0-0.0)	0.0 (0.0 to 0.0)	0.0 (0.0 to 0.0)
Symptomatic cocaine dependence aggregate	1918.5 (1542.1 to 2341.9)	31.1 (28.0 to 34.3)*	14.1 (12.0 to 16.4)*	521.5 (329.0-733.9)	30.6 (27.2 to 34.4)*	14.0 (11.4 to 17.0)*
Amphetamine use disorders	6599.8 (5295.7 to 8024.4)	30.4 (27.0 to 34.2)*	19.2 (16.1 to 22.5)*	874.7 (513.4-1308.7)	30.3 (26.1 to 34.8)*	19.1 (15.5 to 23.0)*
Asymptomatic amphetamine dependence	3613.7 (2722.6 to 4630.7)	30.4 (27.0 to 34.2)*	19.2 (16.1 to 22.5)*	0.0 (0.0-0.0)	0.0 (0.0 to 0.0)	0.0 (0.0 to 0.0)
Symptomatic amphetamine dependence aggregate	2986.1 (2186.5 to 3 893.0)	30.4 (27.0 to 34.2)*	19.2 (16.1 to 22.5)*	874.7 (513.4-1 308.7)	30.3 (26.1 to 34.8)*	19.1 (15.5 to 23.0)*
Cannabis use disorders	19 762.5 (17 982.4 to 21 770.2)	5.3 (4.0 to 6.5)*	-3.8 (-4.7 to -2.9)*	577.2 (371.8-816.2)	5.3 (3.7 to 7.1)*	-3.7 (-5.0 to -2.3)*

(Table 3 continues on next page)

	Prevalence (thousands)			YLDs (thousands)		
	2015	Percentage change in counts between 2005 and 2015	Percentage change in ASR between 2005 and 2015	2015	Percentage change in counts between 2005 and 2015	Percentage change in ASR between 2005 and 2015
(Continued from previous page)						
Asymptomatic cannabis dependence	11 430.9 (10 159.4 to 12 892.9)	5.3 (4.0 to 6.5)*	-3.8 (-4.7 to -2.9)*	0.0 (0.0-0.0)	0.0 (0.0 to 0.0)	0.0 (0.0 to 0.0)
Symptomatic cannabis dependence aggregate	8331.6 (7244.3 to 9593.1)	5.3 (4.0 to 6.5)*	-3.8 (-4.7 to -2.9)*	577.2 (371.8-816.2)	5.3 (3.7 to 7.1)*	-3.7 (-5.0 to -2.3)*
Other drug use disorders	906.4 (550.1-1 312.1)	30.5 (25.7 to 35.6)*	17.2 (12.9 to 21.7)*
Depressive disorders	311 147.6 (300 016.8 to 320 544.3)	18.4 (17.2 to 19.5)*	0.7 (-0.3 to 1.6)	54 255.4 (37 569.9-72 943.3)	18.2 (17.2 to 19.2)*	1.0 (0.5 to 1.5)*
Major depressive disorder	216 047.0 (192 863.4 to 243 319.4)	17.8 (16.6 to 19.0)*	0.8 (0.2 to 1.4)*	44 224.4 (29 672.5-60 297.0)	17.8 (16.6 to 19.0)*	1.1 (0.5 to 1.7)*
Dysthymia	104 106.3 (90 398.1 to 118 968.9)	19.9 (18.4 to 21.5)*	0.6 (-0.3 to 1.4)	10 031.0 (6 605.9-14 267.1)	19.8 (18.3 to 21.5)*	0.7 (-0.2 to 1.6)
Bipolar disorder	44 015.8 (38 150.4 to 50 912.5)	14.9 (14.1 to 15.8)*	0.3 (0.1 to 0.5)*	9004.7 (5501.7-13 388.0)	14.9 (13.9 to 15.9)*	0.5 (-0.0 to 0.9)
Anxiety disorders	267 202.4 (234 064.3 to 306 318.0)	14.9 (13.0 to 16.8)*	0.8 (-0.5 to 2.1)	24 643.0 (16 813.7-33 647.8)	14.8 (12.8 to 16.6)*	1.0 (-0.4 to 2.3)
Eating disorders	6468.4 (6170.6 to 6753.6)	19.1 (16.5 to 21.8)*	9.9 (7.5 to 12.4)*	1386.1 (915.9-1941.2)	19.0 (16.9 to 21.2)*	10.0 (8.1 to 11.9)*
Anorexia nervosa	2912.1 (2357.7 to 3596.2)	12.0 (9.3 to 14.8)*	5.1 (2.6 to 7.6)*	620.5 (411.3-897.7)	12.1 (9.0 to 15.1)*	5.2 (2.3 to 8.0)*
Bulimia nervosa	3629.4 (2937.5 to 4406.8)	25.3 (23.6 to 27.1)*	14.2 (12.5 to 15.8)*	765.6 (490.0-1108.5)	25.3 (23.2 to 27.6)*	14.2 (12.2 to 16.2)*
Autistic spectrum disorders	62 212.4 (58 979.0 to 64 744.3)	12.3 (10.5 to 14.1)*	0.3 (-1.2 to 1.9)	10 051.5 (6740.1-13800.3)	12.3 (11.9 to 12.8)*	0.6 (0.2 to 0.9)*
Autism	24790.1 (20 957.6 to 29 393.1)	12.6 (12.2 to 12.9)*	0.5 (0.3 to 0.7)*	6335.9 (4112.0-8916.1)	12.5 (11.9 to 13.1)*	0.7 (0.2 to 1.2)*
Asperger syndrome and other autistic spectrum disorders	37 245.4 (31 510.5 to 44 883.1)	12.1 (11.8 to 12.4)*	0.2 (0.1 to 0.3)*	3715.6 (2479.6-5424.6)	12.0 (11.5 to 12.5)*	0.3 (-0.1 to 0.7)
Attention-deficit or hyperactivity disorder	51 094.3 (46 162.4 to 57 267.7)	0.3 (-0.4 to 1.0)	-3.6 (-4.2 to -3.0)*	620.1 (370.1-947.7)	0.4 (-0.6 to 1.2)	-3.5 (-4.4 to -2.8)*
Conduct disorder	48 135.5 (39 315.2 to 58 151.6)	0.4 (-0.1 to 1.1)	1.3 (0.9 to 1.6)*	5770.5 (3485.2-8955.7)	0.5 (-0.3 to 1.4)	1.4 (0.7 to 2.0)*
Idiopathic developmental intellectual disability	92 074.0 (52 280.1 to 130 411.2)	10.9 (9.5 to 11.9)*	1.0 (0.0 to 1.8)*	3442.1 (1506.4-5996.9)	10.3 (8.2 to 11.4)*	0.2 (-1.5 to 1.2)
Other mental and substance use disorders	128 178.3 (127 512.5 to 128 877.2)	18.8 (17.9 to 19.7)*	0.2 (-0.6 to 0.9)	9613.4 (6699.7-12 951.9)	18.7 (17.7 to 19.8)*	0.3 (-0.6 to 1.1)
Diabetes, urogenital, blood, and endocrine diseases	2 944 626.8 (2 925 016.2 to 2 968 511.3)	15.5 (14.7 to 16.1)*	-0.3 (-0.9 to 0.3)	66 092.7 (46 781.5-88 865.8)	22.3 (20.8 to 24.0)*	1.2 (0.1 to 2.3)*
Diabetes	435 328.4 (404 736.1 to 468 562.4)	30.6 (28.0 to 33.0)*	4.5 (2.4 to 6.4)*	33 360.8 (23 043.5-45 530.8)	32.5 (29.7 to 35.3)*	5.4 (3.2 to 7.5)*
Uncomplicated diabetes	271 713.6 (241 414.2 to 302 381.7)	27.8 (24.9 to 30.4)*	3.2 (1.1 to 5.3)*	12 599.0 (7979.9-18 394.2)	27.7 (24.8 to 30.3)*	3.3 (1.2 to 5.4)*
Neuropathy and other complications of diabetes aggregate	159 067.7 (134 666.7 to 187 174.0)	35.6 (32.2 to 38.8)*	6.6 (4.1 to 9.0)*	20 459.6 (13 572.0-28 749.6)	35.6 (32.3 to 38.8)*	6.7 (4.2 to 9.0)*
Vision loss due to diabetes aggregate	4547.1 (3823.8 to 5416.0)	34.7 (32.4 to 36.9)*	5.0 (3.6 to 6.3)*	302.3 (210.1-415.5)	36.1 (33.6 to 38.7)*	5.7 (4.1 to 7.4)*
Acute glomerulonephritis	100.5 (92.1 to 110.1)	-0.9 (-4.4 to 1.9)	-14.5 (-17.3 to -12.6)*	5.0 (3.2-7.5)	-0.6 (-4.6 to 2.8)	-14.1 (-17.4 to -11.4)*
Chronic kidney disease	322 510.6 (312 718.2 to 330 351.1)	26.9 (25.8 to 28.0)*	0.9 (-0.0 to 1.8)	8172.8 (6118.6-10 229.3)	23.8 (22.9 to 24.8)*	0.1 (-0.4 to 0.6)
Chronic kidney disease due to diabetes	100 823.9 (86 922.7 to 115 652.3)	27.3 (24.9 to 29.9)*	2.1 (0.5 to 3.7)*	2456.1 (1780.4-3151.2)	23.0 (20.7 to 25.4)*	0.6 (-0.7 to 2.0)

(Table 3 continues on next page)

	Prevalence (thousands)			YLDs (thousands)		
	2015	Percentage change in counts between 2005 and 2015	Percentage change in ASR between 2005 and 2015	2015	Percentage change in counts between 2005 and 2015	Percentage change in ASR between 2005 and 2015
(Continued from previous page)						
Stage 3 chronic kidney disease due to diabetes aggregate	90 937.7 (77 906.5 to 104 617.0)	27.9 (25.4 to 30.5)*	2.3 (0.6 to 4.0)*	119.6 (79.7–175.5)	11.8 (6.5 to 16.7)*	–6.3 (–10.0 to –2.8)*
Stage 4 chronic kidney disease due to diabetes aggregate	6441.3 (5521.3 to 7401.3)	21.6 (19.1 to 24.2)*	0.9 (–0.4 to 2.2)	787.7 (541.8–1 106.1)	20.2 (17.6 to 22.7)*	0.4 (–1.0 to 1.8)
Stage 5 chronic kidney disease untreated due to diabetes	2694.4 (2331.9 to 3091.5)	25.6 (23.0 to 28.2)*	1.3 (–0.0 to 2.8)	1359.3 (920.5–1791.4)	25.3 (22.4 to 28.4)*	1.4 (–0.3 to 3.2)
End-stage chronic kidney disease due to diabetes aggregate	750.5 (643.3 to 864.2)	19.8 (16.4 to 23.4)*	–3.0 (–5.2 to –0.7)*	189.5 (130.5–252.0)	26.6 (22.3 to 30.9)*	0.7 (–2.1 to 3.7)
Chronic kidney disease due to hypertension	78 962.3 (67 846.7 to 91 021.2)	26.0 (23.6 to 28.6)*	0.2 (–1.4 to 1.9)	1508.0 (1108.7–1938.1)	28.5 (25.8 to 31.5)*	1.2 (–0.6 to 3.3)
Stage 3 chronic kidney disease due to hypertension aggregate	72 772.4 (61 662.5 to 84 498.2)	25.9 (23.3 to 28.6)*	0.2 (–1.6 to 2.0)	80.8 (54.2–120.1)	22.1 (17.5 to 26.6)*	–4.4 (–8.0 to –0.9)*
Stage 4 chronic kidney disease due to hypertension aggregate	4102.2 (3503.5 to 4772.6)	25.9 (22.4 to 29.5)*	0.7 (–1.6 to 2.8)	489.8 (341.4–696.7)	25.2 (22.1 to 28.7)*	0.5 (–1.5 to 2.5)
Stage 5 chronic kidney disease untreated due to hypertension	1652.2 (1404.5 to 1913.7)	30.7 (27.5 to 34.2)*	1.8 (–0.5 to 4.1)	816.8 (565.2–1060.3)	30.5 (27.0 to 34.5)*	2.0 (–0.5 to 4.7)
End-stage chronic kidney disease due to hypertension aggregate	435.5 (363.3 to 509.1)	27.5 (22.0 to 33.6)*	0.6 (–3.1 to 4.9)	120.6 (83.1–162.3)	33.2 (27.4 to 39.8)*	3.8 (–0.3 to 8.9)
Chronic kidney disease due to glomerulonephritis	67 348.7 (58 198.9 to 77 439.5)	28.9 (25.8 to 32.3)*	1.1 (–1.1 to 3.6)	1951.4 (1432.2–2479.2)	22.2 (20.1 to 24.4)*	–0.7 (–2.1 to 0.7)
Stage 3 chronic kidney disease due to glomerulonephritis aggregate	59 790.8 (50 844.6 to 69 692.7)	29.9 (26.4 to 33.6)*	1.3 (–1.1 to 4.1)	111.3 (73.5–165.2)	17.1 (13.1 to 20.7)*	–5.6 (–8.8 to –2.7)*
Stage 4 chronic kidney disease due to glomerulonephritis aggregate	4872.4 (4297.2 to 5506.6)	21.3 (18.8 to 23.9)*	–0.5 (–2.2 to 1.6)	602.5 (418.4–845.6)	20.2 (17.7 to 22.9)*	–0.5 (–2.3 to 1.5)
Stage 5 chronic kidney disease untreated due to glomerulonephritis	2228.1 (1895.7 to 2596.9)	23.6 (21.3 to 25.9)*	–0.6 (–2.0 to 0.7)	1122.6 (773.7–1461.3)	23.5 (20.5 to 26.3)*	–0.4 (–2.4 to 1.5)
End-stage chronic kidney disease due to glomerulonephritis aggregate	457.4 (377.1 to 543.0)	20.5 (16.7 to 24.5)*	–2.6 (–5.1 to –0.0)*	115.0 (80.2–152.9)	26.0 (21.1 to 31.2)*	0.4 (–3.2 to 4.1)
Chronic kidney disease due to other causes	94 553.5 (81 141.8 to 109 371.3)	26.2 (24.2 to 28.3)*	0.1 (–1.1 to 1.5)	2257.2 (1663.4–2892.2)	23.0 (21.1 to 25.1)*	–0.5 (–1.6 to 0.6)
Stage 3 chronic kidney disease due to other causes aggregate	85 459.8 (72 471.6 to 99 605.4)	26.6 (24.6 to 28.8)*	0.1 (–1.2 to 1.6)*	132.2 (88.0–194.5)	17.6 (14.0 to 20.8)*	–6.2 (–8.8 to –3.8)*
Stage 4 chronic kidney disease due to other causes aggregate	6058.4 (5257.9 to 6938.4)	21.3 (18.9 to 23.5)*	–0.2 (–1.1 to 0.8)	739.7 (513.5–1 029.6)	19.4 (17.0 to 21.6)*	–1.0 (–2.2 to 0.2)
Stage 5 chronic kidney disease untreated due to other causes	2464.6 (2115.1 to 2835.9)	25.7 (23.8 to 27.5)*	0.1 (–0.9 to 1.1)	1233.0 (855.9–1604.1)	25.5 (23.0 to 28.0)*	0.3 (–1.1 to 1.8)

(Table 3 continues on next page)

	Prevalence (thousands)			YLDs (thousands)		
	2015	Percentage change in counts between 2005 and 2015	Percentage change in ASR between 2005 and 2015	2015	Percentage change in counts between 2005 and 2015	Percentage change in ASR between 2005 and 2015
(Continued from previous page)						
End-stage chronic kidney disease due to other causes aggregate	570.6 (485.6 to 668.1)	21.6 (18.0 to 25.0)*	-2.2 (-4.2 to -0.2)*	152.4 (107.3-200.8)	26.3 (21.8 to 30.7)*	0.1 (-3.0 to 3.2)
Urinary diseases and male infertility	436 081.7 (425 577.7 to 446 450.5)	24.2 (22.8 to 25.4)*	0.9 (-0.1 to 1.9)	4220.9 (2699.9-6045.2)	28.9 (26.8 to 31.0)*	-0.7 (-2.1 to 0.7)
Interstitial nephritis and urinary tract infections	2901.2 (2831.9 to 2980.4)	19.3 (19.0 to 19.7)*	3.1 (2.9 to 3.3)	96.6 (59.9-144.0)	19.2 (17.5 to 20.9)*	3.3 (1.8 to 4.7)*
Urolithiasis	319 600.2 (291 519.2 to 349 966.1)	22.8 (20.8 to 24.9)*	0.9 (-0.7 to 2.5)	89.7 (60.9-122.5)	23.4 (20.0 to 27.0)*	2.8 (0.1 to 5.5)*
Benign prostatic hyperplasia	104 625.3 (90 729.8 to 118 244.0)	29.7 (27.5 to 32.0)*	-1.3 (-2.8 to 0.2)	3792.7 (2423.8-5437.3)	29.8 (27.6 to 32.2)*	-1.3 (-2.8 to 0.3)
Male infertility	29 033.2 (23 972.5 to 34 766.8)	22.6 (19.3 to 25.6)*	9.6 (6.8 to 12.1)*	173.9 (70.1-365.3)*	22.7 (19.3 to 25.9)*	9.8 (6.9 to 12.5)*
Other urinary diseases	67.9 (45.5-96.5)	17.2 (9.1 to 27.2)*	-0.6 (-7.3 to 7.7)*
Gynaecological diseases	794 303.2 (786 497.9 to 801 169.7)	13.8 (13.2 to 14.5)*	-1.8 (-2.4 to -1.2)*	10 001.2 (6807.0-14 312.3)	10.7 (9.4 to 11.9)*	-3.3 (-4.3 to -2.5)*
Uterine fibroids	151 115.0 (144 146.9 to 158 477.5)	19.2 (18.8 to 19.5)*	-0.4 (-0.6 to -0.2)*	2383.5 (1450.2-3834.4)	11.1 (8.6 to 13.2)*	-5.8 (-7.7 to -4.2)*
Uterine fibroids cases aggregate	108 335.0 (101 618.4 to 115 233.6)	23.0 (22.3 to 23.7)*	1.7 (1.2 to 2.1)*	645.9 (314.9-1193.3)	26.1 (25.0 to 27.3)*	3.4 (2.7 to 4.3)*
Mild abdominal pain with anaemia due to uterine fibroids aggregate	42 780.0 (41 634.5 to 43 881.4)	10.5 (9.5 to 11.5)*	-5.5 (-6.3 to -4.6)*	1737.5 (1125.7-2625.4)	6.4 (4.5 to 8.1)*	-8.9 (-10.5 to -7.4)*
Polycystic ovarian syndrome	60 106.4 (46 139.3 to 75 963.4)	10.7 (10.1 to 11.4)*	-0.5 (-0.9 to -0.0)*	532.6 (238.9-1 044.6)	10.8 (9.9 to 11.5)*	-0.5 (-1.1 to 0.0)
Polycystic ovarian syndrome cases aggregate	47 116.8 (35 880.1 to 60 919.9)	10.4 (9.8 to 11.1)*	-0.8 (-1.2 to -0.3)*	408.0 (181.1-797.7)	10.5 (9.7 to 11.3)*	-0.7 (-1.2 to -0.1)*
Hirsutism and infertility due to polycystic ovarian syndrome aggregate	12 989.6 (8 635.6 to 18 975.6)	11.9 (10.9 to 12.7)*	0.4 (-0.4 to 1.1)	124.6 (47.2-278.0)	11.5 (9.6 to 12.9)*	0.1 (-1.5 to 1.3)
Female infertility	61 300.1 (45 889.9 to 79 647.1)	24.6 (19.2 to 30.0)*	11.2 (6.5 to 15.8)*	344.5 (134.3-746.7)	25.0 (19.7 to 30.0)*	11.7 (7.2 to 16.2)*
Endometriosis	10 758.2 (9 165.5 to 12 485.9)	11.6 (10.9 to 12.2)*	-2.6 (-3.0 to -2.2)*	996.4 (647.1-1 384.7)	11.7 (10.4 to 12.9)*	-2.4 (-3.5 to -1.5)*
Endometriosis cases aggregate	10 167.5 (8 652.4 to 11 828.8)	11.5 (10.9 to 12.2)*	-2.7 (-3.1 to -2.3)*	937.3 (610.7-1 312.6)	11.6 (10.3 to 12.9)*	-2.5 (-3.6 to -1.6)*
Abdominal pain and infertility due to endometriosis aggregate	590.7 (415.2 to 787.2)	12.3 (11.2 to 13.3)*	-1.0 (-1.8 to -0.3)*	59.1 (34.2-90.8)	12.3 (9.5 to 15.2)*	-0.9 (-3.4 to 1.5)
Genital prolapse	161 679.1 (142 334.7 to 182 566.5)	17.7 (15.2 to 20.0)*	-6.6 (-8.5 to -4.8)*	503.0 (242.5-904.3)	17.8 (15.4 to 20.3)*	-6.5 (-8.4 to -4.6)*
Premenstrual syndrome	430 696.9 (410 840.7 to 450 494.4)	10.1 (7.9 to 11.8)*	-2.1 (-3.9 to -0.7)*	3621.6 (2249.5-5428.5)	10.2 (8.0 to 12.0)*	-2.0 (-3.8 to -0.4)*
Other gynaecological diseases	45 018.8 (44 057.3 to 46 031.2)	8.0 (7.3 to 8.6)*	-4.2 (-4.8 to -3.7)*	1619.8 (1129.8-2210.0)	6.2 (4.9 to 7.6)*	-5.7 (-6.8 to -4.6)*
Other gynaecological diseases cases aggregate	23 908.1 (22 975.7 to 24 918.0)	13.8 (13.5 to 14.0)*	-0.4 (-0.5 to -0.2)*	936.8 (635.4-1 305.9)	13.8 (12.6 to 15.1)*	-0.2 (-1.2 to 0.8)
Anaemia due to other gynaecological diseases aggregate	21 110.8 (20 900.6 to 21 328.2)	2.1 (0.9 to 3.4)*	-8.2 (-9.3 to -7.1)*	682.9 (461.0-978.0)	-2.7 (-4.6 to -0.7)*	-12.3 (-14.0 to -10.6)*

(Table 3 continues on next page)

	Prevalence (thousands)			YLDs (thousands)		
	2015	Percentage change in counts between 2005 and 2015	Percentage change in ASR between 2005 and 2015	2015	Percentage change in counts between 2005 and 2015	Percentage change in ASR between 2005 and 2015
(Continued from previous page)						
Haemoglobinopathies and haemolytic anaemias	1 571 610.0 (1 545 204.5 to 1 605 099.8)	11.1 (9.5 to 12.6)*	-1.1 (-2.5 to 0.3)	8221.5 (5528.6-11 766.6)	4.3 (2.9 to 5.5)*	-4.9 (-6.0 to -3.9)*
Thalassaemias	439.0 (405.7 to 496.3)	1.4 (-0.1 to 3.4)	-6.5 (-7.8 to -4.6)*	31.3 (21.2-44.4)	1.0 (-3.8 to 6.1)	-7.0 (-11.4 to -2.5)*
β-thalassaemia major cases aggregate	229.2 (222.7 to 238.2)	4.4 (3.6 to 5.2)*	-4.2 (-4.9 to -3.4)*	16.1 (11.0-22.9)	4.1 (-2.5 to 11.5)	-4.7 (-10.6 to 1.9)
Haemoglobin E or β-thalassaemia cases aggregate	67.4 (60.1 to 75.2)	5.4 (2.9 to 8.1)*	-2.5 (-4.8 to 0.1)	5.1 (3.4-7.3)	1.8 (-10.0 to 14.2)	-5.9 (-16.7 to 4.9)
Haemoglobin H disease cases aggregate	139.0 (107.5 to 196.4)	-5.3 (-10.0 to 0.8)	-11.8 (-16.2 to -6.3)*	9.7 (6.4-14.1)	-4.9 (-12.3 to 4.9)	-11.4 (-18.2 to -2.4)*
Heart failure due to thalassaemias aggregate	3.4 (3.2 to 3.6)	26.3 (24.3 to 28.4)*	0.1 (-1.1 to 1.4)*	0.4 (0.3-0.6)	26.4 (16.0 to 38.3)*	0.6 (-9.3 to 11.8)
Thalassaemias trait	279 451.4 (272 818.9 to 287 357.4)	10.5 (10.1 to 11.0)*	-2.1 (-2.5 to -1.7)*	3922.2 (2607.9-5653.6)	6.1 (4.8 to 7.4)*	-3.6 (-4.6 to -2.6)*
β-thalassaemia trait cases aggregate	226 029.6 (220 111.5 to 233 332.0)	10.1 (9.6 to 10.5)*	-2.5 (-3.0 to -2.1)*	3661.7 (2437.6-5270.8)	5.9 (4.7 to 7.2)*	-3.8 (-4.9 to -2.8)*
Haemoglobin E trait cases aggregate	53 421.8 (51 122.3 to 55 882.5)	12.6 (11.5 to 13.7)*	-0.1 (-1.0 to 0.8)	260.5 (174.7-382.8)	8.4 (5.2 to 11.3)*	-0.5 (-3.2 to 2.1)
Sickle-cell disorders	4449.9 (4293.7 to 4600.7)	8.2 (5.2 to 11.9)*	2.8 (-0.2 to 6.2)	371.4 (259.0-518.0)	7.7 (3.3 to 12.3)*	2.3 (-1.8 to 6.6)
Homozygous sickle-cell and severe sickle-cell/β-thalassaemia cases aggregate	4035.1 (3876.0 to 4173.7)	6.5 (3.3 to 10.3)*	1.2 (-1.8 to 4.8)	339.4 (237.7-472.1)	6.5 (2.0 to 11.4)*	1.3 (-2.9 to 5.9)
Haemoglobin SC disease cases aggregate	397.6 (368.2 to 427.4)	29.6 (18.3 to 45.6)*	21.5 (11.0 to 36.5)*	30.1 (21.1-42.1)	22.9 (9.6 to 37.4)*	15.2 (2.8 to 28.9)*
Mild sickle-cell/β-thalassaemia cases aggregate	17.2 (15.9 to 18.6)	16.1 (10.1 to 21.8)*	6.4 (0.8 to 11.5)*	1.9 (1.3-2.6)	13.2 (4.7 to 21.9)*	2.7 (-4.9 to 10.5)
Sickle-cell trait	404 565.9 (381 223.4 to 448 154.6)	19.4 (18.3 to 20.3)*	7.5 (6.5 to 8.3)*	1720.3 (1156.5-2459.0)	10.8 (7.3 to 13.4)*	1.4 (-1.8 to 3.6)
G6PD deficiency	247 073.7 (209 306.7 to 286 712.7)	6.9 (1.4 to 11.9)*	-4.4 (-9.3 to 0.1)	28.4 (19.3-39.6)	1.8 (-2.1 to 6.0)	-7.7 (-11.2 to -4.0)*
G6PD cases aggregate	247 068.5 (209 301.6 to 286 707.7)	6.9 (1.4 to 11.9)*	-4.4 (-9.3 to 0.1)	27.7 (18.9-38.7)	1.2 (-2.8 to 5.5)	-8.1 (-11.6 to -4.3)*
Heart failure due to G6PD deficiency aggregate	5.2 (4.7 to 5.7)	38.1 (36.6 to 39.8)*	7.9 (6.8 to 9.1)*	0.6 (0.4-0.9)	38.1 (34.6 to 41.5)*	7.8 (5.0 to 10.6)*
G6PD trait	728 549.2 (676 734.8 to 781 533.8)	9.8 (7.9 to 11.4)*	-2.7 (-4.3 to -1.2)*	28.0 (19.3-38.9)	4.3 (0.1 to 8.8)*	-5.3 (-9.0 to -1.2)*
Other haemoglobinopathies and haemolytic anaemias	74 385.1 (73 897.6 to 74 861.6)	1.8 (0.9 to 2.7)*	-9.0 (-9.7 to -8.2)*	2119.8 (1423.9-3029.9)	-3.8 (-5.5 to -2.2)*	-12.3 (-13.8 to -11.0)*
Other haemoglobinopathies and haemolytic anaemias cases aggregate	74 273.3 (73 786.0 to 74 751.1)	1.8 (0.9 to 2.7)*	-9.0 (-9.8 to -8.2)*	2067.4 (1384.3-2963.9)	-4.1 (-5.8 to -2.4)*	-12.6 (-14.1 to -11.2)*

(Table 3 continues on next page)

(Continued from previous page)

	Prevalence (thousands)			YLDs (thousands)		
	2015	Percentage change in counts between 2005 and 2015	Percentage change in ASR between 2005 and 2015	2015	Percentage change in counts between 2005 and 2015	Percentage change in ASR between 2005 and 2015
Heart failure due to other haemoglobinopathies and haemolytic anaemias aggregate	111.8 (103.8 to 119.6)	34.9 (33.9 to 36.0)*	2.0 (1.3 to 2.7)*	13.1 (8.9–18.2)	34.9 (31.2 to 38.7)*	2.1 (–0.8 to 5.0)
Endocrine, metabolic, blood, and immune disorders	66 128.9 (65 597.1 to 66 663.8)	4.0 (2.2 to 5.9)*	–7.2 (–8.7 to –5.5)*	2110.5 (1431.5–2993.7)	1.3 (–1.1 to 3.8)	–8.3 (–10.4 to –6.1)*
Endocrine metabolic blood and immune disorders cases aggregate	7577.2 (7382.6 to 7766.6)	19.8 (19.0 to 20.6)*	–0.6 (–1.2 to 0.0)	292.4 (200.2–401.8)	19.5 (18.1 to 21.0)*	–0.5 (–1.6 to 0.6)
Anaemia due to endocrine metabolic blood and immune disorders aggregate	58 430.4 (57 936.0–58 946.7)	2.2 (0.3–4.3)*	–8.0 (–9.7–6.2)*	1804.4 (1216.0–2582.3)	–1.3 (–3.9–1.5)	–9.5 (–11.9–7.0)*
	58 430.4 (57 936.0 to 58 946.7)	2.2 (0.3 to 4.3)*	–8.0 (–9.7 to –6.2)*	1804.4 (1216.0–2582.3)	–1.3 (–3.9 to 1.5)	–9.5 (–11.9 to –7.0)*
Heart failure due to endocrine metabolic blood and immune disorders aggregate	121.4 (113.3 to 130.0)	30.7 (29.6 to 31.9)*	–1.9 (–2.6 to –1.2)*	13.6 (9.4–19.1)	30.7 (27.4 to 34.0)*	–1.7 (–4.3 to 0.8)
Musculoskeletal disorders	1304 100.4 (1 288 602.8 to 1 316 641.4)	20.7 (20.2 to 21.2)*	–0.7 (–1.1 to –0.3)*	146 783.8 (106 764.7–194 473.5)	20.5 (19.6 to 21.5)*	–0.7 (–1.3 to –0.0)*
Rheumatoid arthritis	24 491.2 (22 552.0 to 26 750.7)	23.8 (21.1 to 26.7)*	0.6 (–1.5 to 2.9)	5777.8 (4016.1–7769.6)	23.6 (20.9 to 26.7)*	0.7 (–1.4 to 3.1)*
Osteoarthritis	237 368.6 (230 335.9 to 244 648.1)	32.9 (31.9 to 33.8)*	2.2 (1.6 to 2.9)*	12 886.2 (8 999.7–17 540.0)	34.8 (33.6 to 36.0)*	3.9 (3.0 to 4.8)*
Osteoarthritis of the hip cases aggregate	35 629.2 (32 482.6 to 38 970.1)	33.5 (32.4 to 34.6)*	1.8 (1.0 to 2.6)*	1776.2 (1224.9–2477.0)	35.8 (34.4 to 37.3)*	3.7 (2.6 to 5.0)*
Osteoarthritis of the knee cases aggregate	201 739.4 (195 205.3 to 208 276.6)	32.7 (31.7 to 33.9)*	2.3 (1.5 to 3.1)*	11 110.0 (7742.1–15 123.2)	34.6 (33.3 to 35.9)*	3.9 (3.0 to 4.9)*
Low back and neck pain	820 689.8 (803 467.4 to 837 808.9)	18.7 (17.9 to 19.4)*	–2.0 (–2.6 to –1.4)*	94 941.5 (67 825.3–128 035.0)	18.6 (17.6 to 19.6)*	–2.1 (–2.7 to –1.4)*
Low back pain	539 907.4 (521 448.6 to 559 556.0)	17.3 (16.5 to 18.2)*	–2.8 (–3.4 to –2.2)*	60 074.8 (42 721.9–82 343.7)	17.2 (16.4 to 18.1)*	–2.6 (–3.2 to –2.0)*
Neck pain	358 006.6 (313 408.4 to 409 411.0)	21.1 (19.0 to 23.3)*	–1.1 (–2.4 to 0.0)	34 866.7 (23 362.1–47 636.9)	21.0 (18.9 to 23.2)*	–1.1 (–2.3 to 0.1)
Gout	42 214.2 (37 688.2 to 47 495.5)	26.4 (25.2 to 27.7)*	0.5 (0.1 to 1.0)*	1342.8 (910.0–1843.8)	26.3 (24.6 to 27.9)*	0.6 (–0.3 to 1.5)
Asymptomatic gout	37 712.9 (33 598.4 to 42 467.0)	26.4 (25.2 to 27.7)*	0.5 (0.1 to 1.0)*	0.0 (0.0–0.0)	0.0 (0.0 to 0.0)	0.0 (0.0 to 0.0)
Other musculoskeletal disorders	342 067.7 (305 430.7 to 385 146.7)	20.7 (17.5 to 24.0)*	1.2 (–1.2 to 3.7)	31 835.4 (21 489.1–44 268.4)	20.5 (17.2 to 23.7)*	1.3 (–1.1 to 3.8)
Other non-communicable diseases	5316 342.1 (5 283 701.8 to 5 355 928.3)	14.6 (14.3 to 14.8)*	0.4 (0.2 to 0.6)*	139 001.8 (95 459.0–197 704.3)	20.4 (19.5 to 21.5)*	1.2 (0.6 to 2.0)*
Congenital anomalies	95 706.4 (88 133.4 to 102 483.4)	22.5 (18.5 to 26.3)*	9.3 (5.8 to 12.8)*	8621.0 (5389.1–12 950.2)	28.5 (20.0 to 37.6)*	14.7 (7.1 to 22.7)*
Neural tube defects	1449.6 (988.4 to 2091.8)	17.7 (14.5 to 20.3)*	7.2 (4.3 to 9.7)*	499.1 (285.3–823.7)	18.4 (14.1 to 22.2)*	8.1 (4.1 to 11.6)*
Congenital heart anomalies	48 869.1 (34 325.7 to 72 829.2)	29.8 (25.3 to 34.1)*	15.4 (11.4 to 19.2)*	1688.2 (659.4–3116.9)	29.7 (25.6 to 33.7)*	15.6 (11.7 to 19.1)*
Less severe heart anomalies cases aggregate	44 599.5 (29 800.6 to 68 498.5)	29.4 (24.5 to 34.0)*	15.2 (10.7 to 19.1)*	1431.0 (532.7–2747.7)	29.3 (24.2 to 34.0)*	15.2 (10.7 to 19.2)*

(Table 3 continues on next page)

	Prevalence (thousands)			YLDs (thousands)		
	2015	Percentage change in counts between 2005 and 2015	Percentage change in ASR between 2005 and 2015	2015	Percentage change in counts between 2005 and 2015	Percentage change in ASR between 2005 and 2015
(Continued from previous page)						
Severe congenital heart anomalies	4008.6 (2671.1 to 6011.2)	33.7 (27.4 to 39.0)*	18.5 (12.9 to 23.3)*	235.0 (93.2–467.8)	33.5 (27.4 to 38.7)*	18.5 (13.2 to 23.2)*
Critical congenital heart anomalies	155.8 (90.3 to 252.4)	42.0 (27.5 to 52.9)*	28.6 (15.5 to 38.5)*	9.2 (3.2–18.7)	42.0 (26.4 to 54.2)*	28.9 (14.9 to 40.1)*
Heart failure due to congenital heart anomalies aggregate	105.2 (99.2 to 111.4)	11.6 (10.9 to 12.4)*	–1.0 (–1.5 to –0.5)*	12.9 (8.8–18.2)	11.6 (8.3 to 15.1)*	–1.0 (–3.9 to 2.0)
Cleft lip and cleft palate	6882.6 (4029.2 to 11 483.5)	19.5 (7.7 to 27.5)*	6.6 (–3.9 to 13.9)	79.6 (39.5–143.1)	12.2 (2.6 to 19.9)*	0.9 (–7.9 to 7.7)
Down syndrome	5361.8 (3424.6 to 8192.1)	17.9 (13.7 to 21.8)*	6.4 (2.3 to 10.0)*	507.0 (286.2–837.2)	20.3 (15.8 to 24.2)*	6.5 (2.2 to 10.2)*
Turner syndrome	372.4 (174.6 to 677.1)	11.9 (9.8 to 13.9)*	0.1 (–1.7 to 1.9)	6.6 (2.4–14.4)	11.4 (8.7 to 14.0)*	–0.1 (–2.4 to 2.2)
Klinefelter syndrome	220.5 (115.5 to 395.5)	12.8 (10.9 to 14.2)*	0.5 (–1.2 to 1.9)	1.3 (0.5–3.1)	13.5 (10.6 to 15.6)*	0.2 (–2.2 to 2.0)
Other chromosomal abnormalities	4728.6 (2628.9 to 8303.2)	16.2 (11.8 to 19.8)*	4.3 (0.1 to 7.7)*	454.1 (233.5–852.1)	18.5 (13.6 to 22.3)*	4.2 (–0.5 to 7.9)
Other congenital anomalies	32363.0 (17 788.1 to 59 394.1)	14.5 (12.3 to 16.8)*	2.0 (–0.1 to 4.1)	5385.0 (3209.4–8562.1)	31.3 (18.8 to 45.9)*	17.1 (6.1 to 30.3)*
Skin and subcutaneous diseases	2239493.4 (2 222 716.3 to 2 258 252.8)	12.5 (12.1 to 12.8)*	0.7 (0.4 to 1.1)*	44 896.0 (28 943.3–67 159.6)	11.7 (11.0 to 12.3)*	0.4 (0.1 to 0.7)*
Dermatitis	245290.6 (227 283.4 to 262 752.2)	14.0 (13.2 to 14.8)*	0.3 (0.1 to 0.7)*	8788.0 (5963.4–12 273.5)	13.6 (12.9 to 14.4)*	1.6 (1.0 to 2.2)*
Eczema cases aggregate	85585.6 (75 115.2 to 97 031.7)	13.0 (12.4 to 13.6)*	3.3 (2.8 to 3.8)*	5239.2 (3515.1 to 7348.6)	13.0 (12.2 to 13.9)*	3.4 (2.7 to 4.2)*
Contact dermatitis cases aggregate	98015.1 (85 097.0 to 110 396.7)	15.2 (13.3 to 16.9)*	–1.0 (–1.2 to –0.9)*	2632.0 (1625.8–3852.4)	15.0 (13.1 to 16.8)*	–0.9 (–1.4 to –0.5)*
Seborrhoeic dermatitis cases aggregate	61690.0 (54 642.8 to 68 788.8)	13.3 (12.0 to 14.7)*	–1.3 (–1.7 to –1.0)*	916.8 (521.6–1454.1)	13.2 (11.8 to 14.7)*	–1.3 (–1.8 to –0.8)*
Psoriasis	79699.7 (76 690.9 to 82 804.5)	17.6 (17.0 to 18.3)*	0.4 (–0.1 to 0.9)	6438.3 (4495.6–8734.5)	17.5 (16.6 to 18.4)*	0.6 (–0.0 to 1.2)
Cellulitis	959.9 (887.6 to 1033.6)	24.0 (22.1 to 25.7)*	5.8 (4.3 to 7.2)*	69.0 (45.0–97.8)	23.6 (20.7 to 26.1)*	5.8 (3.5 to 8.1)*
Pyoderma	5812.7 (5581.7 to 6019.0)	15.5 (14.7 to 16.3)*	1.4 (0.7 to 2.1)*	32.7 (13.2–68.4)	15.4 (14.0 to 16.8)*	1.5 (0.3 to 2.7)*
Scabies	204151.7 (177 533.7 to 237 466.2)	6.6 (4.0 to 9.5)*	–2.6 (–4.5 to –0.6)*	5268.9 (2966.9–8605.6)	6.6 (3.8 to 9.5)*	–2.5 (–4.5 to –0.5)*
Fungal skin diseases	492372.6 (448 950.9 to 538 232.5)	13.3 (12.5 to 14.1)*	1.6 (1.3 to 1.9)*	2783.3 (1105.6–5905.3)	13.3 (12.4 to 14.1)*	1.7 (1.4 to 2.0)*
Viral skin diseases	174843.1 (165 156.3 to 185 072.0)	8.5 (8.0 to 9.0)*	–1.1 (–1.4 to –0.8)*	5396.9 (3417.1–7959.9)	8.4 (7.9 to 9.0)*	–1.0 (–1.4 to –0.6)*
Molluscum contagiosum cases aggregate	40464.2 (35 685.6 to 46 233.4)	5.2 (4.7 to 5.8)*	–0.9 (–1.3 to –0.5)*	1263.0 (783.0–1932.3)	5.3 (4.6 to 6.1)*	–0.8 (–1.5 to –0.2)*
Viral warts cases aggregate	134378.9 (125 683.9 to 143 028.0)	9.5 (8.9 to 10.1)*	–1.1 (–1.4 to –0.7)*	4133.9 (2628.4–6153.4)	9.4 (8.8 to 10.1)*	–1.0 (–1.5 to –0.6)*
Acne vulgaris	632741.0 (595 241.9 to 671 248.9)	4.6 (3.5 to 5.6)*	0.8 (–0.1 to 1.7)	6854.0 (3275.1–12 713.9)	4.6 (3.5 to 5.6)*	0.8 (–0.1 to 1.8)
Alopecia areata	20594.9 (19 607.8 to 21 617.0)	14.0 (13.5 to 14.4)*	–1.1 (–1.3 to –1.0)*	695.6 (428.5–1035.6)	13.9 (13.1 to 14.7)*	–1.0 (–1.7 to –0.4)*
Pruritus	69582.5 (62 063.2 to 78 152.6)	17.6 (15.7 to 19.5)*	0.4 (–0.7 to 1.6)	741.6 (360.0–1348.6)	17.5 (15.6 to 19.4)*	0.5 (–0.7 to 1.7)
Urticaria	67749.8 (58 743.2 to 77 086.4)	10.7 (9.6 to 12.0)*	–0.1 (–0.3 to 0.0)	4115.7 (2567.2–5833.6)	10.7 (9.6 to 12.0)*	–0.0 (–0.5 to 0.4)

(Table 3 continues on next page)

	Prevalence (thousands)			YLDs (thousands)		
	2015	Percentage change in counts between 2005 and 2015	Percentage change in ASR between 2005 and 2015	2015	Percentage change in counts between 2005 and 2015	Percentage change in ASR between 2005 and 2015
(Continued from previous page)						
Decubitus ulcer	1087.5 (991.2 to 1189.7)	35.4 (33.4 to 37.4)*	4.7 (3.3 to 6.0)*	161.3 (112.0-218.7)	34.5 (31.9 to 37.3)*	4.7 (2.6 to 6.9)*
Other skin and subcutaneous diseases	605 036.3 (589 500.0 to 619 675.6)	22.8 (22.1 to 23.4)*	2.4 (1.9 to 2.9)*	3550.5 (1706.2-6514.4)	22.6 (22.0 to 23.3)*	2.5 (2.0 to 2.9)*
Sense organ diseases	1788 125.6 (1 771 367.0 to 1 808 438.8)	23.7 (23.3 to 24.1)*	0.6 (0.3 to 0.9)*	68 515.2 (47 798.2-93 894.8)	25.2 (24.2 to 26.4)*	0.6 (-0.0 to 1.3)
Glaucoma	5954.5 (5077.5 to 6905.8)	39.1 (36.9 to 41.4)*	4.2 (2.8 to 5.5)*	541.3 (370.0-747.9)	39.4 (37.0 to 41.9)*	4.3 (2.6 to 5.8)*
Cataract	59 727.0 (53 633.2 to 66 879.2)	26.9 (25.2 to 28.5)*	-3.2 (-4.5 to -2.0)*	3879.7 (2766.9-5229.2)	25.5 (24.1 to 26.9)*	-4.2 (-5.2 to -3.2)*
Macular degeneration	6188.4 (5250.3 to 7273.2)	48.5 (46.1 to 50.8)*	8.2 (6.9 to 9.8)*	462.4 (327.4-633.0)	47.7 (45.2 to 49.9)*	6.9 (5.3 to 8.6)*
Refraction and accommodation disorders	819 307.4 (789 917.4 to 848 059.2)	21.9 (20.5 to 23.2)*	-0.4 (-1.3 to 0.6)	14 593.8 (9392.9-22 901.1)	21.1 (20.2 to 22.1)*	-0.3 (-0.9 to 0.4)
Age-related and other hearing loss	1 210 055.2 (1 140 224.2 to 1 274 664.8)	28.3 (27.4 to 29.2)*	1.4 (1.0 to 1.8)*	40 596.8 (27 898.4-56 075.0)	26.4 (24.6 to 28.3)*	1.1 (0.0 to 2.1)*
Other vision loss	25 797.4 (22 675.6 to 28 504.6)	27.1 (24.6 to 29.5)*	5.5 (4.2 to 6.7)*	1 756.4 (1248.9-2392.7)	29.9 (27.8 to 32.0)*	5.6 (4.3 to 6.7)*
Other sense organ diseases	267 689.7 (258 379.8 to 277 711.4)	24.0 (23.1 to 24.9)*	0.8 (0.1 to 1.4)*	6684.7 (4185.0-9714.7)	23.8 (22.9 to 24.8)*	0.9 (0.2 to 1.5)*
Acute other sense organ diseases	1343.8 (1303.5 to 1385.9)	14.9 (14.2 to 15.7)*	0.9 (0.4 to 1.4)*	35.4 (21.6-51.6)	14.8 (12.7 to 16.9)*	0.9 (-0.9 to 2.7)
Chronic other sense organ diseases	266 345.9 (257 047.4 to 276 383.0)	24.0 (23.1 to 24.9)*	0.8 (0.1 to 1.4)*	6649.3 (4162.3-9662.4)	23.9 (23.0 to 24.9)*	0.9 (0.2 to 1.5)*
Oral disorders	3 521 901.1 (3 468 088.2 to 3 575 854.4)	14.5 (13.8 to 15.0)*	0.4 (-0.1 to 0.9)	16 969.6 (10 296.5-26 044.5)	22.4 (21.6 to 23.2)*	-0.2 (-0.5 to 0.1)
Deciduous caries	572 694.1 (475 089.8 to 686 991.2)	4.5 (2.3 to 6.1)*	-2.3 (-4.2 to -0.7)*	147.2 (63.0-292.1)	4.1 (1.6 to 5.9)*	-2.7 (-4.9 to -0.8)*
Permanent caries	2 521 197.8 (2 361 418.3 to 2 679 668.7)	14.5 (13.7 to 15.4)*	0.8 (0.3 to 1.4)*	1743.4 (776.7-3320.9)	13.5 (12.6 to 14.4)*	-0.4 (-1.2 to 0.4)
Periodontal diseases	537 506.0 (465 113.9 to 625 888.8)	25.4 (24.1 to 26.5)*	1.1 (0.5 to 1.7)*	3520.7 (1359.3-7253.7)	25.4 (24.1 to 26.5)*	1.2 (0.6 to 1.8)*
Edentulism and severe tooth loss	275 619.1 (264 200.8 to 288 252.3)	27.3 (26.9 to 27.7)*	-0.9 (-1.1 to -0.7)*	7640.3 (5096.6-10 562.3)	27.3 (26.9 to 27.7)*	-0.8 (-1.1 to -0.6)*
Other oral disorders	133 719.0 (127 499.7 to 139 776.2)	15.9 (15.4 to 16.4)*	-0.1 (-0.2 to 0.1)	3918.1 (2432.7-5879.4)	15.8 (15.3 to 16.4)*	0.0 (-0.2 to 0.3)
Injuries	1 019 157.4 (977 405.7 to 1 064 704.2)	16.4 (15.5 to 17.4)*	-3.4 (-4.1 to -2.6)*	41 022.0 (29 290.4-55 288.4)	8.0 (4.6 to 11.1)*	-9.9 (-12.5 to -7.5)*
Transport injuries	115 981.0 (110 575.9 to 123 047.1)	23.1 (22.5 to 23.7)*	-0.0 (-0.4 to 0.4)	6444.8 (4500.6-8733.0)	12.2 (8.7 to 15.8)*	-8.0 (-10.7 to -5.3)*
Road injuries	107 722.4 (102 214.5 to 114 850.0)	23.4 (22.8 to 24.1)*	0.3 (-0.1 to 0.7)	5956.1 (4130.8-8076.3)	13.3 (9.9 to 16.7)*	-7.1 (-9.7 to -4.6)*
Pedestrian road injuries	14 157.8 (12 664.5 to 16 095.9)	22.9 (21.7 to 24.1)*	-0.7 (-1.4 to 0.0)	770.8 (532.9-1060.8)	9.6 (5.5 to 13.6)*	-10.6 (-13.7 to -7.6)*
Cyclist road injuries	18 700.6 (16 545.4 to 20 951.1)	21.4 (19.9 to 22.7)*	-0.4 (-1.2 to 0.4)	968.2 (655.2-1348.5)	9.9 (5.5 to 14.2)*	-9.0 (-12.5 to -5.9)*
Motorcyclist road injuries	26 417.2 (23 496.7 to 29 722.0)	31.7 (30.3 to 33.0)*	7.3 (6.5 to 8.1)*	1397.9 (962.0-1919.8)	17.6 (13.0 to 22.1)*	-3.3 (-7.0 to 0.2)
Motor vehicle road injuries	44 311.4 (40 212.6 to 49 496.3)	18.7 (17.7 to 19.8)*	-4.1 (-4.7 to -3.4)*	2585.4 (1811.2-3556.8)	12.2 (9.6 to 14.8)*	-8.4 (-10.4 to -6.6)*

(Table 3 continues on next page)

	Prevalence (thousands)			YLDs (thousands)		
	2015	Percentage change in counts between 2005 and 2015	Percentage change in ASR between 2005 and 2015	2015	Percentage change in counts between 2005 and 2015	Percentage change in ASR between 2005 and 2015
(Continued from previous page)						
Other road injuries	4135.3 (3603.7 to 4692.7)	40.2 (38.6 to 42.1)*	15.2 (14.4 to 16.1)*	233.7 (158.3-320.9)	32.1 (28.3 to 35.5)*	9.5 (6.8 to 11.9)*
Other transport injuries	8258.5 (7469.0 to 9141.6)	18.8 (17.9 to 19.7)*	-3.2 (-3.6 to -2.7)*	488.7 (348.6-660.5)	0.3 (-4.0 to 5.2)	-17.1 (-20.5 to -13.3)*
Unintentional injuries	790101.8 (759 587.4 to 824 995.0)	14.3 (13.8 to 14.8)*	-5.2 (-5.5 to -4.9)*	30 679.5 (21 602.9-41 953.7)	6.2 (3.1 to 9.0)*	-11.4 (-13.6 to -9.2)*
Falls	225736.9 (207 914.3 to 245 787.2)	25.1 (24.4 to 25.8)*	1.7 (1.3 to 2.1)*	11 770.2 (8257.6-16 166.4)	11.3 (6.7 to 15.7)*	-8.6 (-12.1 to -5.2)*
Drowning	4455.5 (4041.2 to 4919.4)	7.0 (5.8 to 8.1)*	-13.1 (-13.5 to -12.7)*	247.3 (173.4-335.8)	-6.7 (-10.3 to -2.4)*	-23.4 (-26.2 to -20.2)*
Fire, heat, and hot substances	69 666.6 (60 988.1 to 78 550.1)	10.8 (10.1 to 11.7)*	-8.3 (-8.8 to -7.9)*	2269.4 (1571.4-3131.8)	1.9 (-1.8 to 5.5)	-13.5 (-16.1 to -11.2)*
Poisonings	5722.4 (4829.3 to 6854.1)	13.8 (12.7 to 14.9)*	-2.0 (-3.2 to -0.6)*	549.4 (367.8-791.1)	8.8 (6.7 to 10.8)*	-4.5 (-6.6 to -2.6)*
Exposure to mechanical forces	183 582.4 (169 658.6 to 197 859.6)	13.9 (13.0 to 14.7)*	-3.5 (-4.0 to -2.9)*	3580.8 (2485.7-4922.9)	5.9 (2.9 to 8.9)*	-9.9 (-12.1 to -7.6)*
Unintentional firearm injuries	2333.0 (2074.5 to 2625.1)	13.4 (12.4 to 14.4)*	-6.3 (-7.1 to -5.6)*	93.2 (65.4-126.2)	4.1 (0.6 to 7.4)*	-12.6 (-15.0 to -10.1)*
Unintentional suffocation	9759.3 (8142.5 to 11 916.9)	9.7 (5.5 to 13.9)*	-7.7 (-10.3 to -5.1)*	353.7 (245.8-480.3)	11.4 (8.6 to 14.0)*	-6.8 (-8.6 to -5.1)*
Other exposure to mechanical forces	171 490.1 (157 821.0 to 185 364.4)	14.1 (13.2 to 14.9)*	-3.2 (-3.7 to -2.6)*	3133.9 (2172.1-4356.6)	5.4 (2.2 to 8.5)*	-10.1 (-12.6 to -7.7)*
Adverse effects of medical treatment	11 158.6 (8773.5 to 13 610.2)	3.1 (1.3 to 4.9)*	-13.9 (-15.4 to -12.4)*	1487.4 (926.9-2223.0)	3.1 (1.3 to 4.9)*	-13.9 (-15.4 to -12.4)*
Animal contact	34 049.7 (30 774.2 to 37 569.0)	9.4 (8.0 to 10.6)*	-6.6 (-7.5 to -5.9)*	1100.9 (763.8-1486.9)	4.6 (3.0 to 6.2)*	-8.8 (-10.0 to -7.5)*
Venomous animal contact	14 680.7 (13 173.4 to 16 375.1)	13.2 (12.2 to 14.2)*	-3.6 (-4.4 to -2.8)*	820.7 (558.1-1117.5)	5.3 (3.7 to 7.1)*	-7.8 (-9.2 to -6.2)*
Non-venomous animal contact	19 369.1 (16 794.5 to 22 306.5)	6.7 (4.8 to 8.2)*	-8.8 (-10.1 to -7.7)*	280.3 (187.2-410.5)	2.5 (0.2 to 4.7)*	-11.4 (-13.1 to -9.8)*
Foreign body	32 345.9 (28 877.3 to 36 026.4)	15.7 (14.1 to 17.1)*	-1.9 (-2.9 to -1.1)*	1263.3 (899.4-1722.5)	3.7 (0.2 to 7.5)*	-10.9 (-13.6 to -8.1)*
Pulmonary aspiration and foreign body in airway	13 351.2 (11 069.0 to 16 798.7)	15.5 (12.5 to 18.4)*	-1.6 (-3.6 to 0.2)	690.2 (477.9-973.1)	-0.2 (-4.1 to 4.5)	-14.0 (-17.2 to -10.4)*
Foreign body in eyes	1007.2 (444.0 to 1 593.4)	16.3 (15.0 to 18.2)*	1.4 (0.8 to 1.9)*	56.2 (24.0-100.6)	15.2 (13.8 to 16.7)*	0.5 (-0.9 to 1.3)
Foreign body in other body part	17 987.4 (15 874.1 to 20 237.9)	15.9 (14.9 to 17.0)*	-2.3 (-3.0 to -1.6)*	517.0 (364.4-713.3)	8.0 (5.5 to 10.6)*	-7.7 (-9.6 to -5.8)*
Environmental heat and cold exposure	61 942.3 (56 211.9 to 68 761.8)	15.9 (15.4 to 16.5)*	-4.2 (-4.6 to -3.8)*	2657.1 (1861.7-3636.2)	6.4 (3.7 to 9.4)*	-11.0 (-13.0 to -8.9)*
Other unintentional injuries	161 441.6 (147 971.3 to 176 204.3)	4.7 (4.0 to 5.3)*	-13.7 (-14.1 to -13.3)*	5753.6 (3913.1-8024.1)	0.4 (-1.4 to 2.1)	-16.7 (-17.8 to -15.7)*
Self-harm and interpersonal violence	24 083.0 (22 279.7 to 25 744.4)	15.2 (14.6 to 15.7)*	-5.5 (-5.9 to -5.2)*	1077.5 (762.7-1 448.2)	1.9 (-1.6 to 5.8)	-15.1 (-17.8 to -12.1)*
Self-harm	6358.6 (5682.1 to 7154.5)	15.2 (14.5 to 15.8)*	-6.2 (-6.7 to -5.8)*	332.7 (232.4-446.9)	0.9 (-2.1 to 4.5)	-15.9 (-18.4 to -13.2)*
Interpersonal violence	17 724.4 (16 186.6 to 19 268.4)	15.2 (14.5 to 15.9)*	-5.3 (-5.7 to -4.8)*	744.8 (521.9-1007.3)	2.4 (-1.4 to 6.4)	-14.6 (-17.5 to -11.6)*
Assault by firearm	1043.6 (900.8 to 1213.2)	16.1 (14.9 to 17.3)*	-4.9 (-5.8 to -4.2)*	46.5 (32.4-64.9)	4.6 (0.7 to 8.1)*	-13.2 (-16.1 to -10.5)*
Assault by sharp object	4019.5 (3330.4 to 4668.1)	12.5 (11.5 to 13.6)*	-7.0 (-7.7 to -6.3)*	123.4 (84.8-168.7)	-6.4 (-11.4 to -0.9)*	-21.1 (-24.8 to -16.8)*
Assault by other means	12 661.3 (11 425.5 to 14 018.9)	16.0 (15.2 to 16.8)*	-4.7 (-5.3 to -4.3)*	574.8 (404.0-782.2)	4.3 (0.8 to 8.0)*	-13.3 (-15.9 to -10.4)*

(Table 3 continues on next page)

	Prevalence (thousands)			YLDs (thousands)		
	2015	Percentage change in counts between 2005 and 2015	Percentage change in ASR between 2005 and 2015	2015	Percentage change in counts between 2005 and 2015	Percentage change in ASR between 2005 and 2015
(Continued from previous page)						
Forces of nature, war, and legal intervention	88 991.7 (65 131.0 to 116 786.1)	28.8 (18.4 to 40.5)*	11.7 (2.9 to 21.7)*	2820.2 (1921.3 to 3850.1)	23.4 (1.7 to 47.8)*	7.2 (-11.4 to 28.3)
Exposure to forces of nature	25 293.7 (16 193.6 to 37 658.7)	13.5 (-2.1 to 32.2)	-1.3 (-14.5 to 14.1)	796.9 (541.4 to 1091.4)	-27.4 (-43.2 to -6.2)*	-35.9 (-49.6 to -17.8)*
Collective violence and legal intervention	63 697.9 (45 248.7-83 186.4)	36.1 (20.9-54.9)*	18.1 (5.1-34.4)*	2023.3 (1303.8-2903.0)	70.5 (33.0-118.3)*	47.1 (15.3-88.8)*
Impairments
Anaemia	2 359 107.9 (2 349 938.3 to 2 369 065.8)	4.0 (3.5 to 4.5)*	-7.1 (-7.6 to -6.7)*	77 876.7 (52 436.9 to 111 360.0)	-0.7 (-2.0 to 0.4)	-10.2 (-11.2 to -9.2)*
Developmental intellectual disability	152 664.0 (113 511.5 to 190 764.1)	12.6 (12.2 to 13.0)*	2.0 (1.6 to 2.4)*	16 875.1 (12 072.6 to 22 373.1)	15.0 (13.4 to 17.0)*	3.8 (2.2 to 5.7)*
Epilepsy	39 160.5 (34 270.8 to 43 685.8)	15.8 (12.6 to 18.9)*	3.5 (0.5 to 6.4)*	11 770.0 (8 578.8 to 15 276.3)	5.8 (0.5 to 11.0)*	-4.9 (-9.8 to -0.1)*
Guillain-Barré syndrome	35.0 (28.6 to 42.1)	17.0 (14.7 to 19.5)*	-0.2 (-0.9 to 0.4)	10.4 (6.6 to 15.0)	17.0 (14.7 to 19.5)	-0.2 (-0.9 to 0.4)
Hearing loss	1 330 902.9 (1 261 401.0 to 1 397 100.6)	26.0 (25.2 to 26.7)*	0.9 (0.5 to 1.3)*	46 183.1 (31 566.7 to 63 209.9)	23.6 (22.1 to 25.1)*	0.6 (-0.4 to 1.5)
Heart failure	40 048.6 (38 613.2 to 41 418.2)	32.3 (31.6 to 33.1)*	0.3 (-0.2 to 0.8)	6199.4 (4700.7 to 7785.5)	33.2 (31.7 to 34.6)*	1.1 (0.1 to 2.0)*
Infertility	113 113.8 (93 429.4-136 497.5)	21.1 (17.6-24.5)*	8.0 (4.9-10.9)*	752.4 (327.7-1543.8)	19.9 (16.8-22.7)*	7.0 (4.3-9.4)*
Pelvic inflammatory disease	754.1 (652.8 to 878.7)	1.1 (0.1 to 2.1)*	-12.0 (-12.7 to -11.2)*	99.9 (67.8 to 138.6)	1.3 (-0.7 to 3.4)	-11.7 (-13.4 to -10.0)*
Vision loss	939 580.2 (905 009.5 to 974 112.3)	22.4 (21.2 to 23.6)*	-0.4 (-1.3 to 0.4)	24 462.7 (16 954.9 to 34 455.1)	22.4 (21.6 to 23.3)*	-0.2 (-0.8 to 0.3)

Data in parentheses are 95% UIs. *Percentage changes that are statistically significant.

Table 3: Global prevalence and YLDs for 2015, percentage change of counts, and percentage change of age-standardised rates between 2005 and 2015 for all causes, Level 5 sequelae, and nine impairments

disability in Nepal, Bangladesh, and Bhutan, respectively. YLDs due to anxiety were lower than expected in all countries in the region except for Bangladesh.

In the GBD super-region of central Europe, eastern Europe, and central Asia, lower back and neck pain was the leading cause of disability in every geographical region in 2015; Although all countries in eastern and central Europe recorded higher than expected YLD ratios for most of their top ten causes, central Asia mostly had lower than expected or as expected YLD ratios. Sense organ diseases were the second leading causes of disability and depressive disorders were the third leading causes in this super-region.

Sizeable discrepancies occurred for observed and expected YLDs based on SDI throughout north Africa and the Middle East, probably reflecting the uneven achievements in development found in this region. Lower back and neck pain was the primary driver of YLDs in the region. Exceptions to this were Lebanon and Syria, for which war caused the most disability in 2015, and Afghanistan for which iron-deficiency was the leading cause. Depression, sense organ diseases, diabetes ranked second, third, and fourth, respectively, for the

region. Observed YLDs due to diabetes consistently exceeded expected levels, with two countries posting ratios higher than 3.00 (ie, Qatar [3.12], and the United Arab Emirates [3.52]). Iran and Morocco recorded much higher disability due to drug use disorders than expected based on SDI.

In comparison with the rest of the world, ranks of cause-specific disability, and their ratios of observed to expected values, were vastly different in sub-Saharan Africa. Of the 46 countries within the super-regions, nine had lower back and neck pain as the leading cause of YLDs in 2015. In southern Sub-Saharan Africa, HIV/AIDS was the leading cause of disability for all countries. Iron-deficiency anaemia ranked as the leading cause of YLDs for 11 countries in western sub-Saharan Africa. For the remaining countries, lower back and neck pain were primarily the leading cause of YLDs; Liberia, the exception, had onchocerciasis as its leading cause of disability. Malaria and various neglected tropical diseases caused more YLDs than expected in most west African countries. For eastern sub-Saharan Africa, sense organ diseases, iron deficiency, depressive disorders, and lower back and neck pain were leading causes of YLDs

in 2015. In central sub-Saharan Africa, skin diseases were consistently among the leading three-to-four causes of disability in this region. HIV/AIDS resulted in far more YLDs than expected based on SDI, with Equatorial Guinea, and Central African Republic recording ratios of observed versus expected YLDs higher than 2·00.

Discussion

We used 60900 data sources to estimate the incidence and prevalence of 2619 sequelae of 310 causes for 591 geographical regions for the years 1990, 1995, 2000, 2005, 2010, and 2015. After accounting for variation over time and across regions in case definitions, disease assays, survey instruments, and coding practice, and using standardised modelling approaches to deal with missing data, conflicting data, and a range of sampling and non-sampling errors, we characterised the global and national patterns in non-fatal health outcomes. We found that global age-standardised YLDs per capita decreased slightly in the last 25 years from 0·114 (0·085–0·147) in 1990 to 0·110 (0·082–0·141) per capita in 2015, a total decrease of 3·69% (3·12–4·25%) over a generation (appendix pp 714–16). The ageing of the world's population and the general increase in YLDs per capita with age resulted in an increase in global YLDs per capita. Within this overall pattern, 133 out of 310 causes had statistically significant decreases in YLDs per capita between 2005 and 2015 compared with 82 causes with statistically significant increases in YLDs per capita over the same time period. The most important contributors to global YLDs were musculoskeletal disorders (18·5% [16·4–20·9%] of all YLDs in 2015), mental and substance use disorders (18·4% [15·6–21·2%]), and the category of other NCDs (17·9% [15·0–21·7%]), dominated by hearing loss, vision loss, and skin diseases.

Typology of diseases based on YLDs and YLLs

The comparison of trends in age-standardised YLDs and YLLs provides insight into the drivers of differences in rates of increases and decreases for diseases and injuries. Diseases and injuries can be parsed into four groups. The first of these are a small set of disorders including causes such as drug use disorders and skin cancer for which age-standardised rates of both YLDs and YLLs increased at a rate of more than 0·4% per year from 2005 to 2015. A second category of diseases includes those in which YLDs are increasing but YLLs are decreasing, including disorders such as thyroid cancer, cirrhosis, and neonatal encephalopathy. One explanation for this pattern is that risk factors, broadly defined, are causing an increase in incidence, whereas access to effective treatment has improved enough to continue reducing mortality. For a third category of disorders, YLDs are essentially constant, but YLLs are decreasing. This grouping is quite large and includes many of the major causes of disability such as musculoskeletal disorders, various neurological disorders, and a number of cancers. For many infectious causes, decreased are occurring for both YLLs and YLDs but are

much faster for YLLs; this pattern is to be expected if access to effective treatment is decreasing the number of deaths but not reducing prevalence or incidence. The last category includes disorders where decreases are equal, if not slightly higher, for YLDs including ischaemic heart disease, falls, cervical cancer, and other injuries. Examination of differential trends by region or country might provide insights into the role of access to treatment in reducing YLLs and, conversely, the role of increasing or decreasing risks and social determinants in driving changes in disease incidence and prevalence. More detailed exploration of these differential trends is warranted in future work.

Disability and retirement age

As life expectancy has steadily increased in most countries, there have been calls in some high SDI countries to extend retirement ages to reflect these changes in survival.^{28–30} A crucial factor in these debates is whether individuals are living for longer and have higher average levels of functional health at each age or not. In this study, we found that age-standardised YLDs per capita decreased, but rather slowly, over the period 2005 to 2015. Comparison of trends in age-standardised YLDs and YLLs by cause shows that the main causes of YLDs are not decreasing and in some settings might be increasing. Other than a somewhat slower rate of improvement for YLDs at oldest age (≥ 80 years), age discontinuities are not observable in these patterns. Ultimately, the debate on retirement age will hinge on the skill sets required for different types of work, whether work contributes to diminished or improved functional health status, and societal expectations for retirement. Our findings suggest that the burden from mental and substance use disorders, and musculoskeletal disorders, which are frequent causes of early retirement^{31,32} in terms of age-specific prevalence, has not declined much over time; the continued prevalence of these and other disorders associated with increasing age might limit the capacities of an older workforce.

Epidemiological transition

The analysis of YLD rates by Level 2 causes as a function of SDI shows a very different picture than that reported for YLLs. At the lower end of the SDI spectrum, incremental increases in SDI are associated with reductions in both age-standardised YLDs and all-age YLD rates spurred by decreases in disability from neglected tropical diseases and anaemia as well as decreases in tuberculosis, diarrhoea, pneumonia, meningitis, and other infectious diseases. At SDI levels above 0·8 (see methods appendix pp 698–711 for SDI values for each country), age-standardised rates of health loss increase slightly, attributable to higher rates of mental and substance use disorders, musculoskeletal disorders, and neurological disorders. In this phase of the epidemiological transition, this increase in rates, although small, combined with population ageing, results in increases in YLDs per 100 000. The rising average YLD per capita rates have implications for health systems; a larger

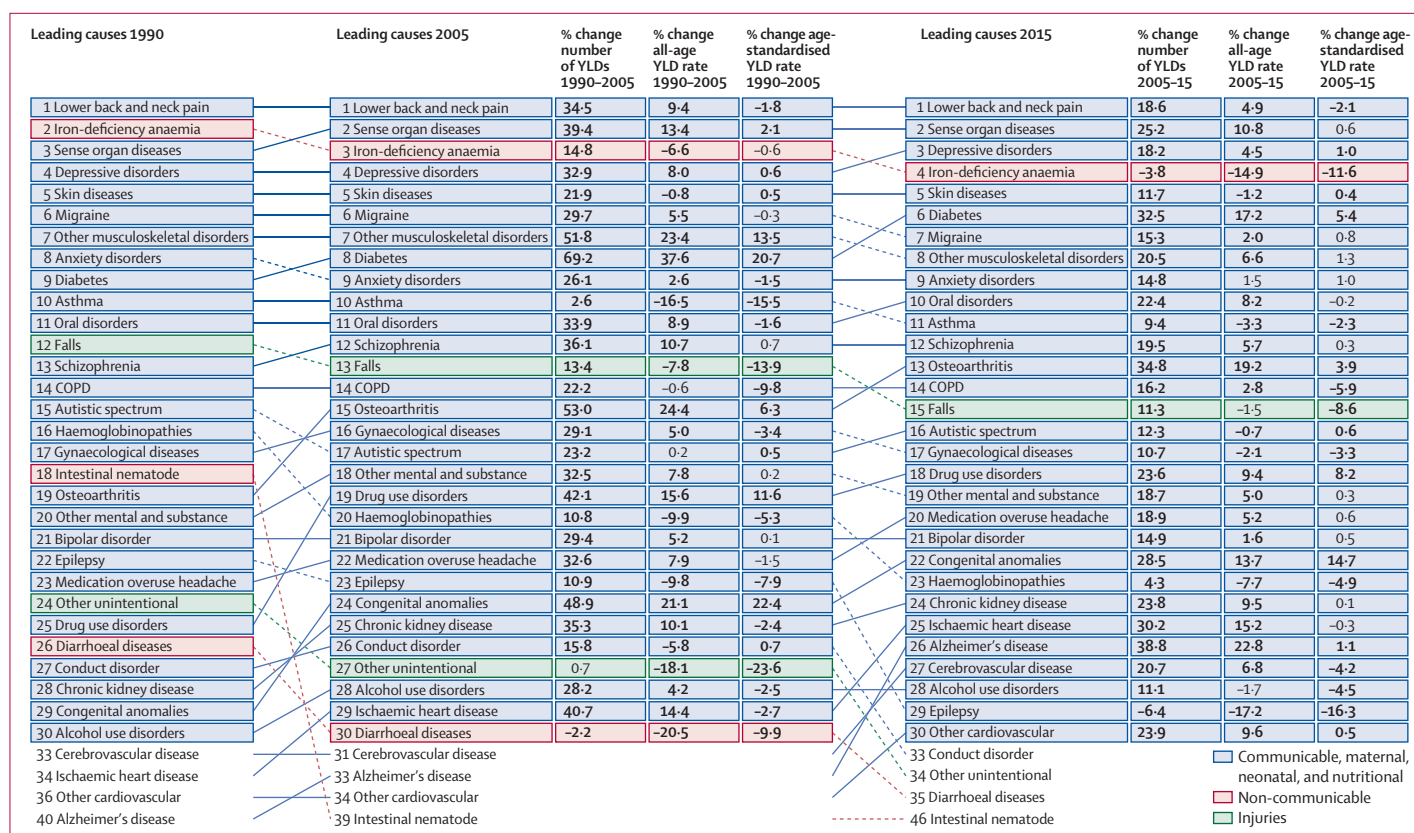


Figure 2: Leading 30 Level 3 causes of global YLDs for both sexes combined, 1990, 2005, and 2015, with percentage change in number of YLDs, and all-age and age-standardised rates
Causes are connected by lines between time periods. For the time period of 1990 to 2005 and for 2005 to 2015, three measures of change are shown: percent change in the number of YLDs, percent change in the all-age YLD rate, and percent change in the age-standardised YLD rate. YLD=years lived with disability. COPD=chronic obstructive pulmonary disease.

fraction of the population is likely to need care for many disorders. Some of these disorders are currently costly to manage.³³ The increase in health-care costs per individual in the population that occurs once an SDI of 0.8 is exceeded is a predictable component of the epidemiological transition; these costs should be anticipated during the health planning processes of countries in that stage of the transition.

Disease-specific issues

Musculoskeletal disorders

Musculoskeletal disorders continue to be a leading cause of disability worldwide and more so when taking into account that additional musculoskeletal burden from long-term sequelae of fractures and dislocations is classified under injuries in GBD. A key component of healthy ageing is to maintain mobility, and a key public health intervention recommended for improving health outcomes for all chronic diseases is physical activity. Painful musculoskeletal disorders increase with age and are a great threat to mobility, compromising health more broadly.³⁴⁻³⁶ Even if cures for musculoskeletal disorders are not yet available, the clinical goal of preventing disability is attainable.^{37,38}

Mental and substance use disorders

Consistent with the findings of earlier GBD studies, GBD 2015 confirms the large contribution of mental and substance use disorders to global disability. For the first time, in GBD 2015 we found a positive association between conflict and depression and anxiety, albeit with wide uncertainty. This uncertainty is due to sparse and low-quality data for these disorders in post-conflict countries. Going forward, we plan to make separate estimates for post-traumatic stress disorder in GBD, which might add considerably more data from conflict settings and show a stronger association. GBD results have provided an evidence base to support global action such as a stated commitment by the World Bank and WHO to make mental health a global development priority³⁹ and the consideration of mental health and substance use disorders in shaping the Sustainable Development Goals.⁴⁰ Cost-effective interventions are able to reduce the burden imposed by mental and substance use disorders, including in low-income and middle-income countries.⁴¹

Increases in deaths attributable to drug use in the USA have resulted in considerable policy and media attention. In our assessment, the ten countries with the

	1	2	3	4	5	6	7	8	9	10
Early neonatal	Iron	NN sepsis	PEM	Haemog	Other inf	Diarrhoea	NN preterm	Congenital	Endocrine	NN enceph
Late neonatal	Iron	PEM	Diarrhoea	Congenital	Haemog	NN preterm	Other nutr	NN enceph	Other inf	Epilepsy
Post neonatal	Iron	Diarrhoea	PEM	Haemog	Skin	Other NTD	Congenital	Other inf	NN preterm	Endocrine
1-4 years	Iron	Skin	PEM	Diarrhoea	Sense	Asthma	Haemog	Other NTD	Congenital	Otitis
5-9 years	Iron	Skin	Asthma	Sense	Haemog	Other NTD	Conduct	Malaria	ASD	Anxiety
10-14 years	Iron	Skin	Conduct	Anxiety	Asthma	Migraine	Sense	Depression	Back & neck	Haemog
15-19 years	Skin	Depression	Iron	Back & neck	Migraine	Anxiety	Sense	Conduct	Other MSK	Asthma
20-24 years	Depression	Back & neck	Skin	Migraine	Iron	Other MSK	Anxiety	Sense	Other mental	Drugs
25-29 years	Back & neck	Depression	Migraine	Skin	Iron	Other MSK	Anxiety	Sense	Drugs	Schiz
30-34 years	Back & neck	Depression	Migraine	Skin	Iron	Sense	Other MSK	Anxiety	Schiz	Gynae
35-39 years	Back & neck	Depression	Migraine	Sense	Other MSK	Skin	Iron	Anxiety	Diabetes	Schiz
40-44 years	Back & neck	Depression	Sense	Migraine	Other MSK	Diabetes	Skin	Iron	Anxiety	Schiz
45-49 years	Back & neck	Depression	Sense	Diabetes	Other MSK	Migraine	Skin	Iron	Anxiety	Schiz
50-54 years	Back & neck	Sense	Depression	Diabetes	Other MSK	Migraine	Skin	Osteoarth	Anxiety	Schiz
55-59 years	Back & neck	Sense	Diabetes	Depression	Other MSK	Migraine	Osteoarth	Skin	Oral	Anxiety
60-64 years	Back & neck	Sense	Diabetes	Depression	Other MSK	Osteoarth	Oral	Skin	Migraine	COPD
65-69 years	Sense	Back & neck	Diabetes	Depression	Other MSK	Osteoarth	Oral	COPD	Skin	IHD
70-74 years	Sense	Back & neck	Diabetes	Depression	Oral	Other MSK	Osteoarth	COPD	IHD	Skin
75-79 years	Sense	Back & neck	Diabetes	Alzheimer's	Depression	Oral	Osteoarth	Other MSK	COPD	IHD
≥80 years	Sense	Alzheimer's	Back & neck	Diabetes	Falls	IHD	Osteoarth	Depression	COPD	Oral

Rate of change 2005-15

-0.19 to -0.03 -0.03 to 0.01 0.01 to 0.06 0.06 to 0.09 0.09 to 0.15
 0.15 to 0.19 0.19 to 0.24 0.24 to 0.29 0.29 to 0.42 0.42 to 0.63

Figure 3: Leading ten Level 3 causes of global age-specific years lived with disability in 2015

Each cause is coloured by the percentage change in age-specific years lived with disability from 2005 to 2015. Alzheimer's=Alzheimer disease and other dementias.

ASD=autism. Back & neck=low back and neck pain. Conduct=conduct disorders. Congenital=congenital anomalies. COPD=chronic obstructive pulmonary disease.

Drugs=drug use disorders. Endocrine=endocrine, metabolic, blood, and immune disorders. Gynae=gynaecological disorders. Haemog=haemoglobinopathies and haemolytic anaemias. IHD=ischaemic heart disease. Iron=iron-deficiency anaemia. NN enceph=neonatal encephalopathy due to birth asphyxia and trauma. NN preterm=neonatal preterm birth complications. NN sepsis=neonatal sepsis and other neonatal infections. Other NTD=other neglected tropical diseases. Oral=oral disorders.

Osteoarth=osteoarthritis. Other inf=other infectious diseases. Other mental=other mental and substance use disorders. Other MSK=other musculoskeletal disorders.

Other nutr=other nutritional deficiencies. PEM=protein-energy malnutrition. Schiz=schizophrenia. Sense=sense organ disease. Skin=skin and subcutaneous diseases.

highest prevalence of opioid dependence in decreasing order in 2015 were Iran, United Arab Emirates, Russia, Morocco, the USA, Australia, Ukraine, Tunisia, Belarus, Canada, and Iraq. Among these countries, we estimated the highest excess mortality from opioid dependence in the eastern European countries at about twice the level of that in the USA, Canada, and the north African and Middle Eastern countries; the lowest rate among these highly prevalent countries was in Australia. Within the USA, prescription opioids have been estimated to account for 37% of drug overdose deaths in 2013.^{42,43} The availability of overdose response treatments in the form of naloxone kits to laypersons has also accelerated.⁴⁴ An alternative approach is opioid substitution treatment to reduce the risk of overdose. The intensity at which countries use harm-reduction strategies such as needle exchange and opioid substitution programmes follows an inverse pattern,⁴⁵ with the most intense programmes in Australia, but very rare use of such strategies in eastern Europe, suggesting that embracing harm reduction is an effective means of reducing drug deaths.

Diabetes

To obtain standard estimates for health loss due to diabetes for GBD 2015 using both fasting plasma glucose (FPG) means and standard deviations as well as diabetes prevalence, we re-extracted all available data from 1990 to 2015. Across studies, we found 20 different case definitions

for diabetes. We also included, wherever possible, studies reporting on mean FPG but not diabetes prevalence, using a regression between mean FPG and diabetes prevalence from studies reporting on both. We have also made the assessment of diabetes prevalence more consistent with cause of death data for diabetes. Our improved efforts at measuring the prevalence of diabetes confirms the increase in global age-standardised incidence, prevalence, and YLDs. Of note, the increase in YLDs at the global scale is greater than the increase in YLLs, which reflects improved access to treatments that lower case fatality. The rise of diabetes prevalence, related to the global increase in body-mass index,⁴⁶ given the costs of treating the disease⁴⁷ and the related increases in cardiovascular risks, poses one of the more important challenges to health systems in the coming years. This is particularly the case in regions with high prevalence of diabetes such as central America, north Africa and the Middle East, and Oceania.

Ezzati and colleagues⁴⁸ and the International Diabetes Federation (IDF)⁴⁹ have estimated diabetes prevalence for many countries; the intra-class correlation coefficient for the estimates from Ezzati and colleagues and the GBD for 2015 is 0.74, and for the IDF estimates is 0.65. These large differences appear to stem from the inclusion of cause of death data in the modelling for GBD and also from the inclusion of self-reported diabetes prevalence in the study by Ezzati and colleagues.⁴⁸ In GBD, these sources were excluded from our analyses because of changing patterns in the prevalence of known and

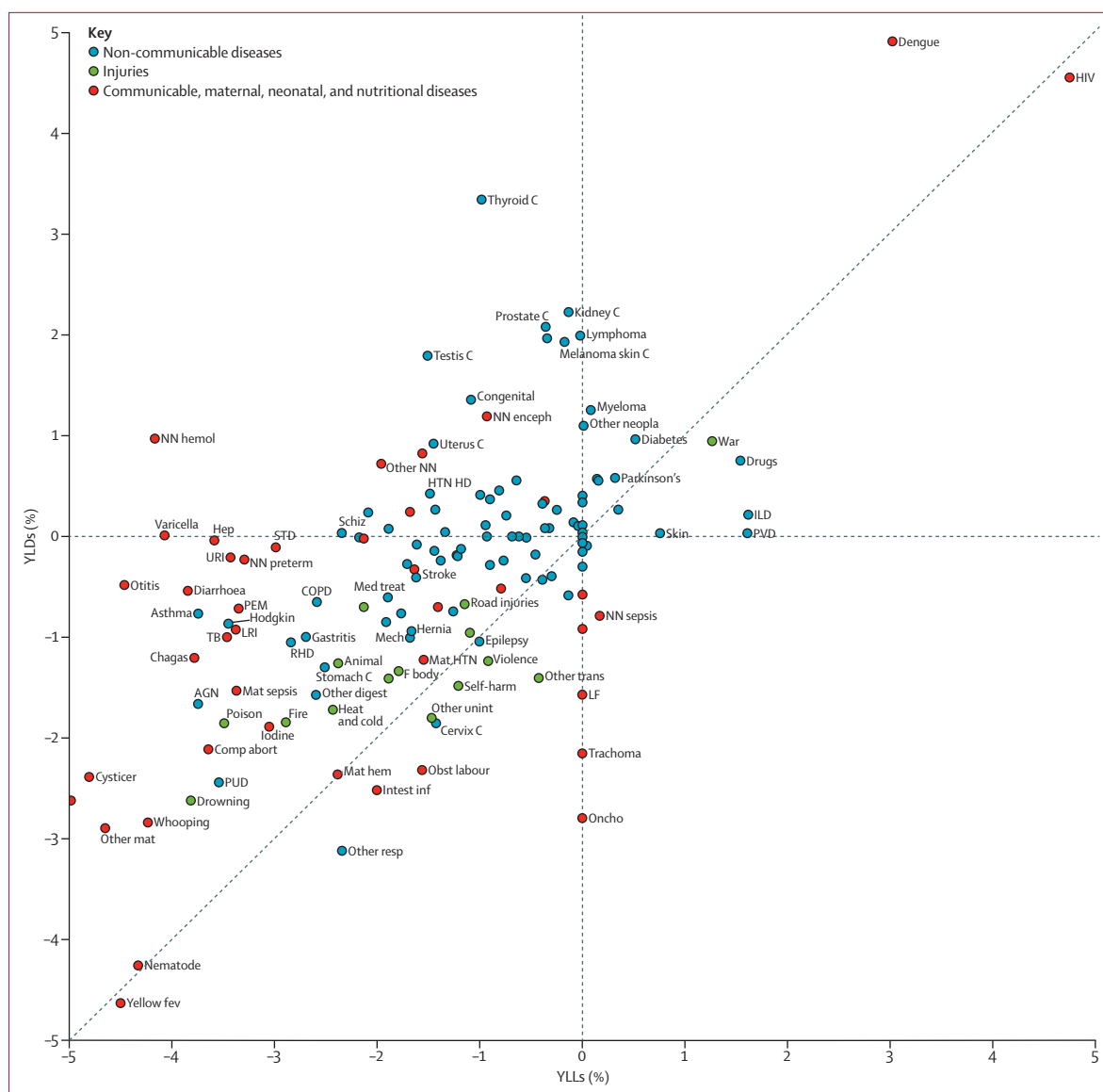


Figure 4: Global annualised rate of change in age-standardised years of life lost (YLLs) and years lived with disability (YLDs) for Level 3 causes between 1990 and 2015

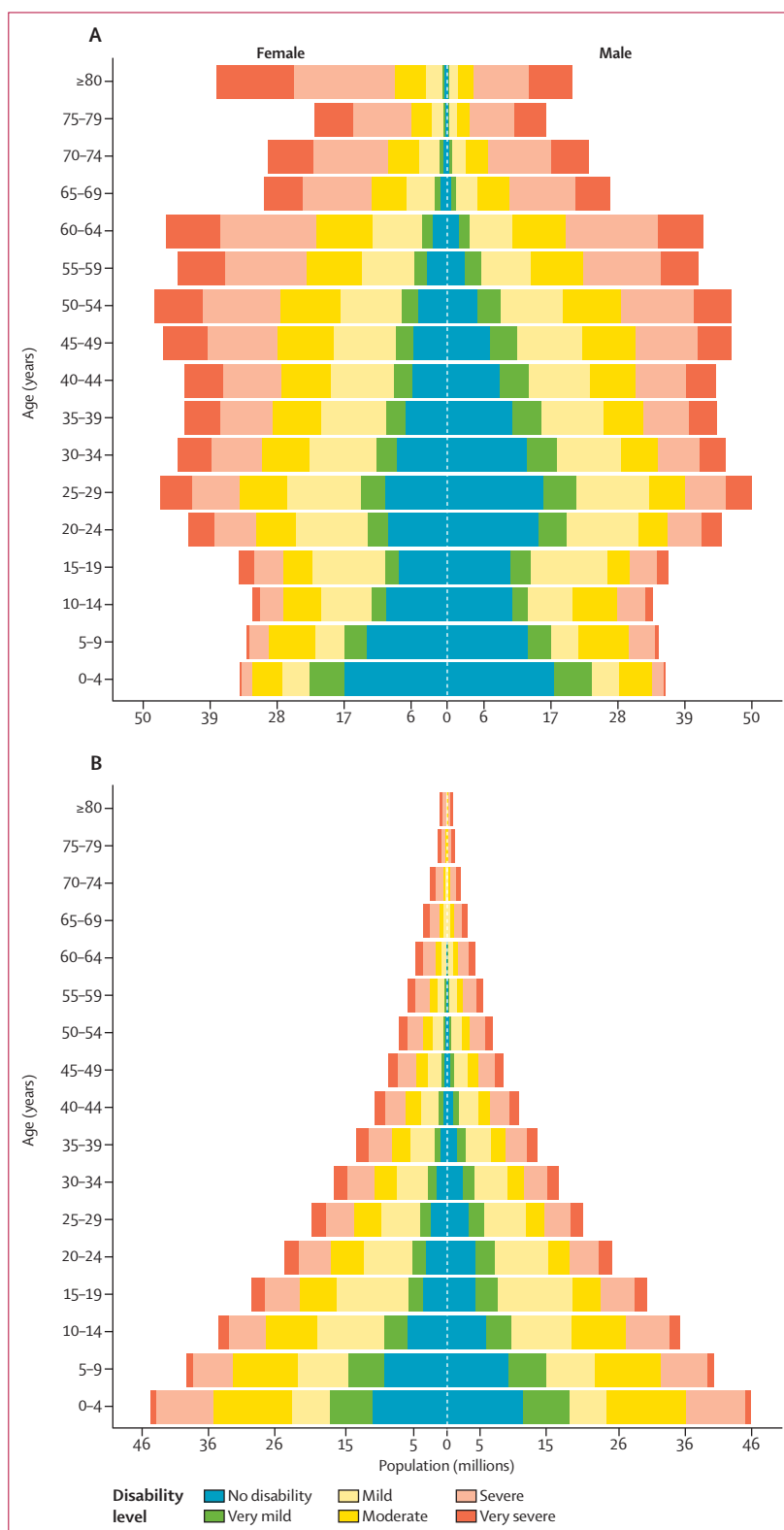
TB=tuberculosis. HIV=HIV/AIDS. Diarrhoea=diarrhoeal diseases. Intest Inf=intestinal infectious diseases. LRI=lower respiratory infections. URI=upper respiratory infections. Otitis=otitis media. Whooping=whooping cough. Varicella=varicella and herpes zoster. Chagas=chagas disease. Cysticer=cysticercosis. LF=lymphatic filariasis. Oncho=onchocerciasis. Trachoma=trachoma. Dengue=dengue. Yellow Fev=yellow fever. Nematode=intestinal nematode infections. Mat hem=maternal haemorrhage. Mat sepsis=maternal sepsis and other maternal infections. Mat HTN=maternal hypertensive disorders. Obst labour=maternal obstructed labour and uterine rupture. Comp abort=maternal abortion, miscarriage, and ectopic pregnancy. Oth mat=other maternal disorders. NN preterm=neonatal preterm birth complications. NN enceph=neonatal encephalopathy due to birth asphyxia and trauma. NN sepsis=neonatal sepsis and other neonatal infections. NN haemol=haemolytic disease and other neonatal disorders. Oth NN=other neonatal disorders. PEM=protein-energy malnutrition. Iodine=iodine deficiency. Oth nutr=other nutritional deficiencies. STD=sexually transmitted diseases excluding HIV. Hep=hepatitis. Stomach C=stomach cancer. Melanoma=malignant skin melanoma. Skin C=non-melanoma skin cancer. Cervix C=cervical cancer. Uterus C=uterine cancer. Prostate C=prostate cancer. Testis C=testicular cancer. Kidney C=kidney cancer. Thyroid C=thyroid cancer. Hodgkin=Hodgkin lymphoma. Lymphoma=non-Hodgkin lymphoma. Myeloma=multiple myeloma. Oth neopla=Other neoplasms. RHD=rheumatic heart disease. Stroke=cerebrovascular disease. HTN HD=hypertensive heart disease. PVD=peripheral vascular disease. COPD=chronic obstructive pulmonary disease. Asthma=asthma. ILD=interstitial lung disease and pulmonary sarcoidosis. Oth resp=other chronic respiratory diseases. PUD=peptic ulcer disease. Gastritis=gastritis and duodenitis. Hernia=inguinal, femoral, and abdominal hernia. Oth digest=other digestive diseases. Parkinson=Parkinson's disease. Schiz=schizophrenia. Drugs=drug use disorders. AGN=acute glomerulonephritis. Congenital=congenital anomalies. Skin=skin and subcutaneous diseases. Road inj=road injuries. Oth trans=other transport injuries. Drown=drowning. Fire=fire, heat, and hot substances. Poison=poisonings. Mech=exposure to mechanical forces. Med treat=adverse effects of medical treatment. Animal=animal contact. F body=foreign body. Heat & cold=environmental heat and cold exposure. Oth unint=other unintentional injuries. Violence=interpersonal violence. War=collective violence and legal intervention.

unknown diabetes in surveys that vary with geography, making it difficult to crosswalk these studies.

Dementia

Brayne and colleagues^{50,51} reported that age-specific prevalence of Alzheimer's was decreasing in the UK. Of the four studies in the USA with similar measurements over time that were reviewed, one showed a decrease, whereas no change in prevalence was seen in the remaining studies.^{52,53} Our assessment of age-standardised prevalence showed that Alzheimer's and other dementias decreased slowly in the UK, by 3.69% (2.53–4.85%) from 2005 to 2015, but remained constant in the rest of the world. The decrease might be due to reductions in vascular dementias rather than reductions in Alzheimer's and other dementias. Our assessment also suggests that the number of individuals with Alzheimer's disease increased from 21.7 million (18.9–24.8 million) worldwide in 1990 to 46.0 million (40.2–52.7 million) in 2015. Although it is useful to know that age-standardised rates might be starting to decrease, if only slightly, the rapid increase in the absolute number of cases points to the major challenge that dementia presents to societies with increasing life expectancy. This number is similar to the most recent estimates from the World Alzheimer Report 2015 that estimated 46.8 million people living with dementia in 2015. However, there is less agreement about new cases per year: The World Alzheimer Report estimated 9.9 million new cases of dementia per year in 2015 compared with 6.44 million (5.52–7.45 million) new cases in 2015 estimated in GBD 2015. The World Alzheimer Report separately analysed prevalence and incidence whereas we use DisMod-MR 2.1 to produce internally consistent prevalence and incidence estimates. We observed a fundamental disagreement in the underlying incidence and prevalence data in many countries and decided to exclude incidence data from our modelling approach, putting greater trust in prevalence studies than incidence studies, arguing that the point of incidence in dementia is more difficult to establish than prevalence when the diagnosis over time becomes more established.

Figure 5: Population pyramids with the number of individuals, by age and sex, grouped by severity of their disability weight (DW) for all comorbid conditions combined into no disability, very mild disability (DW 0.0–0.01), mild disability (DW 0.01–0.05), moderate disability (DW 0.05–0.1), severe disability (DW 0.1–0.3), and very severe disability (DW >0.3) for geographies of high (A) and low (B) quintiles of Socio-demographic Index in 2015
Disability weights are combined multiplicatively as $1 - (1 - DW_1)(1 - DW_2) \dots (1 - DW_n)$ for n comorbid sequelae. Socio-demographic Index (SDI) is calculated for each geography as a function of lag-dependent income per capita, average educational attainment in the population aged over 15 years, and the total fertility rate. SDI units are interpretable; a zero represents the lowest level of income per capita, educational attainment, and highest total fertility rate (TFR) observed from 1980 to 2015 and a one represents the highest income per capita, educational attainment, and lowest TFR observed in the same period. Cutoffs on the SDI scale for the quintiles have been selected based on examination of the entire distribution of geographies between 1980 and 2015.



Hepatitis C

The availability of new medical treatments for chronic hepatitis C has driven considerable interest in the prevalence data for this disease. Available treatments are

highly effective but costly.⁵⁴ Some programmes have been launched in Egypt and other lower-SDI countries that offer treatment at lower cost. Good data on the number of courses of treatment that have been delivered are not widely available. Given the number of individuals with chronic infection and the potential to be cured, tracking the fraction of people treated each year should be undertaken. Even in high-income countries such as the USA, the fraction of those treated among those who would benefit from treatment is not yet available.

Malaria

Our assessment of malaria prevalence and incidence in high burden sub-Saharan Africa was based on the Malaria Atlas Project spatial analysis of prevalence surveys done across Africa between 2000 and 2015.^{55,56} The dynamic map of malaria prevalence was updated for GBD 2015 and extended back to 1980. Children younger than 5 years in sub-Saharan Africa showed a remarkable 33·5% (24·8–46·0%) decrease in incidence from the 2005 peak of 0·78 (0·56–0·96) cases per person per year to 0·52 (0·33–0·71) in 2015. Yet there were 81·7 million (52·0–111·7 million) incident cases in this age group in 2015. The decrease in children was more rapid than that recorded for older age groups. In adolescents and adults, incidence dropped by 30·0% (17·2–39·2%) and 21·1% (15·3–26·9%), respectively, and resulted in 58·7 million (37·7–91·5 million) and 53·3 million (40·7–67·5 million) incident cases, respectively, in 2015. The observed decreases in malaria incidence underscore the remarkable effect of the scale-up of antimalarial interventions⁵⁷ across Africa in the past decade.⁵⁵ Sustaining the levels of these interventions is crucial to continue to reduce malaria morbidity on the continent, avoid resurgence, and improve on these levels that are vital to global aspirations for malaria eradication.⁵⁸ Overall, we estimate there were 286·9 million (219·7–377·3 million) cases of malaria worldwide in 2015, whereas the World Malaria Report (WMR) 2015 reports 214 million (149–303 million) cases in 2015.⁵⁹ Of these cases, the WMR estimates that 88% of cases occurred in the WHO African region (188 million) whereas we estimate 189·4 million (151·7–239·4 million; 66·1%) cases for GBD 2015. Outside of Africa, the WMR reports 26 million cases versus our estimate of 97·3 million (50·2–181·8 million), but both sets of estimates feature identical regional rankings among the six WHO regions with malaria. Increasing convergence in estimates is expected as dynamic maps of

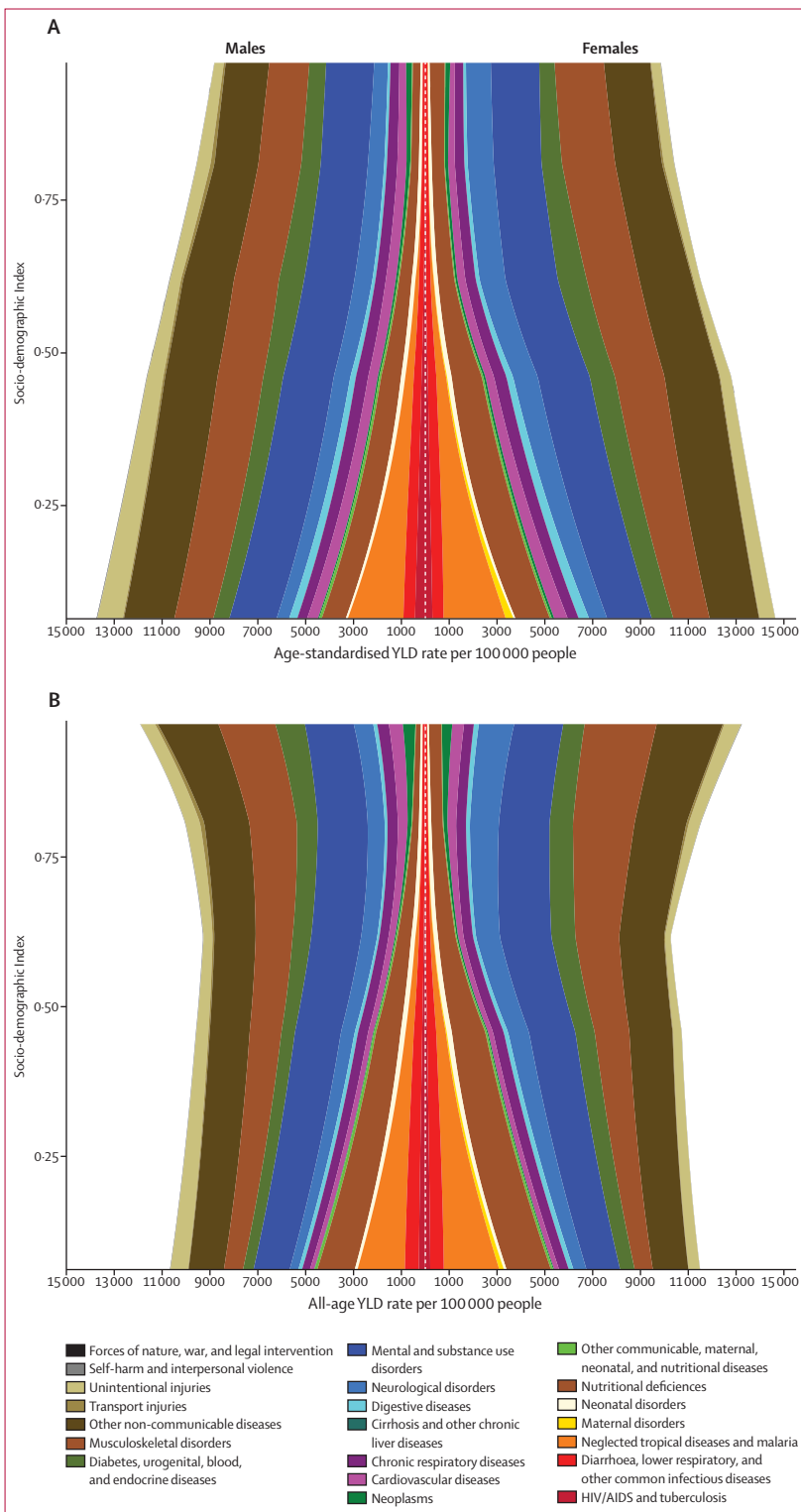


Figure 6: Expected relationship between age-standardised years lived with disability (YLD) rates per 100 000 people for the 21 GBD Level 2 causes and SDI (A) and the expected relationship between all-age years lived with disability (YLD) rates per 100 000 people for the 21 GBD Level 2 causes and SDI (B) by sex. These stacked curves represent the average relationship between SDI and each cause of YLDs observed across all geographies over the time period 1990 to 2015. In each figure, the y-axis spans from lowest SDI up to highest SDI. To the left of the midline are male rates, and the female rates are to the right; higher rates are further from the midline. SDI=Socio-demographic Index.

	1	2	3	4	5	6	7	8	9	10
Global	Back & neck (0.96)	Sense (1.0)	Depression (0.93)	Iron (1.01)	Skin (0.93)	Diabetes (0.98)	Migraine (0.93)	Other MSK (1.29)	Anxiety (0.93)	Oral (0.86)
High income	Back & neck (1.11)	Sense (0.94)	Depression (1.11)	Skin (0.92)	Diabetes (1.29)	Migraine (1.04)	Other MSK (1.27)	Anxiety (1.32)	Oral (0.95)	Iron (0.92)
High-income North America	Back & neck (1.08)	Depression (1.31)	Diabetes (2.3)	Sense (0.89)	Other MSK (1.83)	Skin (0.94)	Anxiety (1.54)	Migraine (0.97)	Drugs (2.94)	Iron (0.9)
Canada	Back & neck (1.31)	Sense (0.91)	Depression (0.98)	Skin (1.02)	Other MSK (1.62)	Diabetes (1.57)	Migraine (1.13)	Iron (1.17)	Anxiety (1.23)	Oral (0.95)
Greenland	Back & neck (1.1)	Depression (0.94)	Other MSK (2.18)	Skin (1.03)	Sense (0.68)	Migraine (1.03)	Anxiety (1.28)	Iron (0.99)	Diabetes (0.72)	Falls (1.17)
USA	Back & neck (1.05)	Depression (1.34)	Diabetes (2.39)	Sense (0.89)	Other MSK (1.86)	Skin (0.93)	Anxiety (1.57)	Migraine (0.95)	Drugs (3.02)	Iron (0.87)
Australasia	Back & neck (1.12)	Depression (1.3)	Sense (0.84)	Other MSK (1.93)	Skin (0.93)	Anxiety (1.72)	Migraine (1.17)	Diabetes (1.27)	Asthma (1.59)	Drugs (2.91)
Australia	Back & neck (1.12)	Depression (1.33)	Other MSK (2.0)	Sense (0.84)	Skin (0.93)	Migraine (1.21)	Anxiety (1.72)	Diabetes (1.22)	Drugs (3.05)	Asthma (1.58)
New Zealand	Back & neck (1.15)	Depression (1.14)	Sense (0.85)	Anxiety (1.77)	Skin (0.92)	Diabetes (1.51)	Other MSK (1.53)	Migraine (1.0)	Asthma (1.63)	Oral (1.09)
High-income Asia Pacific	Back & neck (0.84)	Sense (1.04)	Depression (0.86)	Skin (0.98)	Diabetes (0.96)	Migraine (0.89)	Oral (0.96)	Other MSK (0.92)	Iron (1.12)	Falls (0.89)
Brunei	Back & neck (0.82)	Depression (0.9)	Skin (0.93)	Sense (1.05)	Diabetes (2.39)	Iron (1.06)	Migraine (0.84)	Falls (1.35)	Anxiety (0.85)	Asthma (1.13)
Japan	Back & neck (0.82)	Sense (1.05)	Depression (0.87)	Skin (1.0)	Oral (0.97)	Diabetes (0.83)	Migraine (0.91)	Other MSK (0.93)	Alzheimer's (1.05)	Iron (1.06)
Singapore	Back & neck (0.6)	Sense (1.01)	Depression (0.95)	Skin (0.93)	Migraine (0.75)	Iron (1.05)	Oral (0.95)	Other MSK (0.83)	Anxiety (0.85)	Falls (0.92)
South Korea	Back & neck (0.9)	Sense (1.02)	Depression (0.83)	Diabetes (1.37)	Skin (0.92)	Migraine (0.83)	Iron (1.28)	Other MSK (0.9)	Oral (0.94)	Falls (1.06)
Western Europe	Back & neck (1.27)	Sense (0.94)	Depression (1.05)	Migraine (1.17)	Skin (0.87)	Anxiety (1.28)	Diabetes (0.88)	Oral (0.99)	Other MSK (0.94)	Falls (1.04)
Andorra	Back & neck (1.41)	Sense (0.91)	Depression (1.12)	Skin (0.88)	Migraine (1.15)	Oral (1.13)	Falls (1.23)	Anxiety (1.24)	Diabetes (0.94)	Iron (0.82)
Austria	Back & neck (1.22)	Sense (0.93)	Depression (1.07)	Migraine (1.29)	Skin (0.88)	Oral (1.23)	Anxiety (1.22)	Diabetes (0.9)	Falls (1.19)	Other MSK (0.86)
Belgium	Back & neck (1.39)	Sense (1.06)	Depression (1.01)	Skin (0.87)	Migraine (1.14)	Diabetes (1.1)	Oral (1.17)	Falls (1.25)	Anxiety (1.17)	Other MSK (0.83)
Cyprus	Back & neck (1.4)	Sense (0.96)	Depression (1.07)	Migraine (1.14)	Skin (0.88)	Diabetes (1.27)	Anxiety (1.22)	Oral (1.09)	Asthma (1.24)	Other MSK (0.79)
Denmark	Back & neck (1.45)	Sense (0.79)	Depression (1.1)	Skin (0.94)	Migraine (1.09)	Other MSK (1.23)	Diabetes (1.12)	Oral (1.07)	Anxiety (1.25)	Iron (0.79)
Finland	Back & neck (1.38)	Sense (0.85)	Depression (1.23)	Skin (0.92)	Falls (1.58)	Migraine (1.14)	Diabetes (1.06)	Oral (1.12)	Asthma (1.35)	Iron (1.0)
France	Back & neck (1.15)	Sense (0.92)	Depression (1.03)	Skin (0.86)	Anxiety (1.54)	Migraine (1.04)	Falls (1.23)	Oral (1.0)	Other MSK (1.02)	Diabetes (0.69)
Germany	Back & neck (1.48)	Sense (0.98)	Depression (1.08)	Migraine (1.23)	Skin (0.86)	Anxiety (1.47)	Diabetes (1.02)	Oral (1.05)	Falls (1.02)	Other MSK (0.82)
Greece	Back & neck (1.28)	Sense (0.95)	Depression (1.21)	Migraine (1.11)	Skin (0.83)	Oral (1.13)	Anxiety (1.19)	Diabetes (0.71)	Falls (0.86)	Alzheimer's (1.01)
Iceland	Back & neck (1.38)	Sense (0.93)	Depression (0.89)	Migraine (1.15)	Skin (0.79)	Anxiety (1.23)	Oral (1.18)	Iron (0.83)	Diabetes (0.81)	Falls (0.96)
Ireland	Back & neck (1.25)	Sense (0.92)	Depression (1.07)	Skin (0.91)	Migraine (1.14)	Anxiety (1.58)	Iron (1.0)	Other MSK (1.09)	Oral (1.17)	Asthma (1.36)
Israel	Back & neck (1.2)	Sense (0.94)	Depression (1.1)	Skin (0.86)	Migraine (1.14)	Iron (1.11)	Diabetes (1.2)	Oral (1.15)	Other MSK (0.95)	War (6977.72)
Italy	Back & neck (1.31)	Sense (1.09)	Depression (1.06)	Migraine (1.39)	Diabetes (1.01)	Skin (0.86)	Anxiety (1.24)	Other MSK (1.0)	Alzheimer's (1.22)	Falls (0.94)
Luxembourg	Back & neck (1.42)	Sense (0.92)	Depression (1.11)	Migraine (1.28)	Skin (0.87)	Diabetes (1.3)	Oral (1.21)	Anxiety (1.23)	Falls (1.15)	Asthma (1.42)
Malta	Back & neck (1.31)	Sense (0.89)	Depression (1.02)	Diabetes (1.02)	Migraine (1.13)	Skin (0.9)	Oral (1.15)	Anxiety (1.18)	Falls (1.13)	Asthma (1.12)
Netherlands	Back & neck (1.5)	Sense (0.9)	Depression (0.99)	Migraine (1.21)	Skin (0.87)	Anxiety (1.59)	Diabetes (1.23)	Oral (1.32)	Other MSK (1.13)	Falls (0.94)
Norway	Back & neck (1.59)	Sense (0.81)	Depression (1.03)	Anxiety (1.87)	Skin (0.85)	Migraine (1.07)	Diabetes (1.22)	Oral (1.15)	Other MSK (0.87)	Falls (1.07)
Portugal	Back & neck (1.34)	Sense (0.83)	Depression (1.09)	Migraine (1.13)	Skin (0.88)	Diabetes (0.78)	Oral (1.09)	Anxiety (1.13)	Other MSK (0.94)	Asthma (1.18)
Spain	Back & neck (0.97)	Sense (0.99)	Depression (1.09)	Migraine (1.15)	Skin (0.86)	Diabetes (0.78)	Iron (1.25)	Falls (1.02)	Anxiety (1.01)	Oral (0.91)
Sweden	Back & neck (1.25)	Sense (0.78)	Depression (1.05)	Diabetes (1.36)	Skin (0.9)	Migraine (0.91)	Anxiety (1.22)	Other MSK (1.02)	Oral (0.9)	Iron (0.89)

(Figure 7 continues on next page)

	1	2	3	4	5	6	7	8	9	10
Switzerland	Back & neck (1.48)	Sense (0.9)	Depression (1.08)	Skin (0.88)	Migraine (1.08)	Oral (1.22)	Falls (1.41)	Anxiety (1.25)	Diabetes (0.89)	Other MSK (0.74)
UK	Back & neck (1.17)	Sense (0.79)	Depression (0.96)	Skin (0.88)	Migraine (1.1)	Asthma (1.86)	Other MSK (1.18)	Anxiety (1.07)	Oral (0.91)	Iron (0.92)
England	Back & neck (1.17)	Sense (0.79)	Depression (0.95)	Skin (0.87)	Migraine (1.04)	Other MSK (1.22)	Asthma (1.79)	Anxiety (1.08)	Oral (0.91)	Iron (0.95)
Northern Ireland	Back & neck (1.11)	Sense (0.73)	Depression (0.97)	Skin (0.88)	Migraine (1.03)	Asthma (1.41)	Falls (1.21)	Oral (0.91)	Anxiety (0.89)	Diabetes (0.63)
Scotland	Back & neck (1.23)	Migraine (1.75)	Sense (0.79)	Depression (0.97)	Asthma (2.35)	Skin (0.89)	Diabetes (1.07)	Other MSK (1.02)	Falls (1.21)	Anxiety (1.12)
Wales	Back & neck (1.15)	Sense (0.79)	Depression (1.07)	Asthma (2.59)	Skin (0.9)	Migraine (1.03)	Diabetes (0.95)	Other MSK (1.08)	Falls (1.07)	Anxiety (1.09)
Southern Latin America	Back & neck (0.88)	Depression (1.07)	Sense (0.84)	Skin (0.93)	Anxiety (1.53)	Migraine (0.93)	Iron (1.02)	Diabetes (0.83)	Other MSK (1.07)	Asthma (1.03)
Argentina	Back & neck (0.87)	Depression (1.07)	Sense (0.84)	Skin (0.92)	Anxiety (1.52)	Iron (1.12)	Migraine (0.91)	Diabetes (0.82)	Other MSK (1.09)	Asthma (1.01)
Chile	Back & neck (0.88)	Depression (1.1)	Sense (0.86)	Skin (0.96)	Anxiety (1.57)	Migraine (0.95)	Diabetes (0.9)	Other MSK (1.05)	Oral (0.98)	Asthma (1.08)
Uruguay	Back & neck (0.89)	Sense (0.8)	Depression (1.03)	Skin (0.93)	Anxiety (1.49)	Iron (1.09)	Migraine (0.94)	Diabetes (0.6)	Asthma (1.11)	Other MSK (0.92)
Central Europe, eastern Europe, and central Asia	Back & neck (1.2)	Sense (1.16)	Depression (1.1)	Skin (0.89)	Iron (1.45)	Migraine (0.99)	Diabetes (1.0)	Oral (0.99)	Anxiety (0.81)	Osteoarthritis (1.29)
Eastern Europe	Back & neck (1.21)	Sense (1.25)	Depression (1.17)	Iron (1.81)	Skin (0.89)	Migraine (1.01)	Diabetes (0.93)	Oral (0.96)	Drugs (3.26)	Osteoarthritis (1.33)
Belarus	Back & neck (1.19)	Sense (1.18)	Depression (1.15)	Skin (1.02)	Migraine (1.02)	Diabetes (0.97)	Falls (1.22)	Oral (1.13)	Oral (0.89)	Drugs (2.84)
Estonia	Back & neck (1.16)	Sense (1.08)	Depression (1.25)	Skin (0.88)	Migraine (1.03)	Diabetes (1.05)	Iron (1.16)	Oral (0.93)	Osteoarthritis (1.38)	Alcohol (5.28)
Latvia	Back & neck (1.24)	Sense (1.22)	Depression (0.97)	Diabetes (1.32)	Skin (0.85)	Migraine (1.03)	Iron (1.18)	Oral (0.89)	Osteoarthritis (1.35)	Anxiety (0.77)
Lithuania	Back & neck (1.23)	Sense (1.17)	Depression (1.16)	Skin (0.86)	Diabetes (1.08)	Migraine (1.04)	Iron (1.18)	Oral (0.95)	Osteoarthritis (1.32)	IHD (1.94)
Moldova	Back & neck (1.2)	Sense (0.96)	Depression (1.0)	Skin (0.95)	Migraine (1.02)	Iron (1.24)	Diabetes (0.67)	Oral (0.94)	Anxiety (0.7)	Alcohol (3.29)
Russia	Back & neck (1.2)	Sense (1.3)	Depression (1.14)	Iron (2.22)	Skin (0.88)	Migraine (1.0)	Diabetes (0.95)	Drugs (3.39)	Oral (0.98)	Alcohol (6.05)
Ukraine	Back & neck (1.26)	Sense (1.17)	Depression (1.29)	Skin (0.89)	Migraine (1.03)	Diabetes (0.86)	Oral (0.92)	Osteoarthritis (1.24)	Drugs (3.26)	IHD (1.76)
Central Europe	Back & neck (1.29)	Sense (1.1)	Depression (1.06)	Diabetes (1.14)	Skin (0.9)	Migraine (0.98)	Oral (1.01)	Anxiety (0.93)	Osteoarthritis (1.29)	Iron (0.91)
Albania	Back & neck (1.37)	Sense (0.94)	Depression (0.98)	Skin (0.83)	Iron (1.25)	Migraine (0.98)	Anxiety (0.88)	Oral (0.98)	Diabetes (0.51)	Osteoarthritis (0.91)
Bosnia	Back & neck (1.16)	Sense (0.98)	Depression (0.98)	Diabetes (1.01)	Skin (0.9)	Migraine (0.98)	War (1887.24)	Oral (1.06)	Iron (1.01)	Anxiety (0.88)
Bulgaria	Back & neck (1.28)	Sense (1.04)	Depression (1.06)	Diabetes (1.18)	Skin (0.9)	Migraine (0.98)	Oral (1.01)	Anxiety (0.96)	Osteoarthritis (1.21)	Iron (1.12)
Croatia	Back & neck (1.37)	Sense (0.97)	Depression (1.04)	Skin (0.9)	Diabetes (0.83)	Migraine (0.97)	Oral (1.07)	Iron (1.1)	Osteoarthritis (1.2)	Anxiety (0.92)
Czech Republic	Back & neck (1.38)	Sense (1.09)	Depression (1.13)	Diabetes (1.46)	Skin (0.88)	Migraine (0.98)	Oral (1.04)	Iron (1.05)	Anxiety (0.96)	Osteoarthritis (1.43)
Hungary	Back & neck (1.38)	Sense (1.05)	Depression (1.07)	Diabetes (1.62)	Skin (0.96)	Migraine (0.98)	Oral (0.94)	Anxiety (0.95)	Osteoarthritis (1.35)	Iron (0.85)
Macedonia	Back & neck (1.26)	Sense (0.99)	Depression (1.01)	Diabetes (1.14)	Skin (0.89)	Migraine (0.98)	Oral (0.97)	Anxiety (0.89)	Osteoarthritis (1.21)	Iron (0.58)
Montenegro	Back & neck (1.27)	Sense (1.04)	Depression (1.04)	Diabetes (1.15)	Skin (0.91)	Migraine (0.98)	Oral (1.12)	Iron (0.98)	Anxiety (0.91)	Osteoarthritis (1.26)
Poland	Back & neck (1.21)	Sense (1.27)	Depression (1.07)	Diabetes (1.25)	Skin (0.88)	Migraine (0.97)	Oral (1.03)	Anxiety (0.95)	Osteoarthritis (1.34)	IHD (2.09)
Romania	Back & neck (1.33)	Sense (1.03)	Depression (1.04)	Skin (0.9)	Migraine (0.98)	Diabetes (0.75)	Oral (1.04)	Iron (1.18)	Osteoarthritis (1.28)	Anxiety (0.89)
Serbia	Back & neck (1.31)	Sense (1.0)	Depression (1.02)	Diabetes (1.22)	Skin (0.92)	Migraine (0.98)	Oral (0.98)	Anxiety (0.91)	Osteoarthritis (1.23)	Iron (0.8)
Slovakia	Back & neck (1.34)	Sense (1.06)	Depression (1.08)	Skin (0.89)	Diabetes (1.25)	Migraine (0.97)	Iron (1.09)	Anxiety (0.94)	Oral (0.85)	Osteoarthritis (1.35)
Slovenia	Back & neck (1.35)	Sense (0.99)	Depression (1.09)	Skin (0.9)	Diabetes (1.07)	Migraine (0.98)	Oral (1.16)	Iron (1.18)	Osteoarthritis (1.35)	Anxiety (0.95)
Central Asia	Back & neck (1.01)	Sense (0.98)	Depression (0.97)	Iron (1.36)	Skin (0.87)	Migraine (0.98)	Diabetes (0.98)	Anxiety (0.8)	Oral (1.06)	Asthma (0.69)
Armenia	Back & neck (1.0)	Sense (1.03)	Depression (1.01)	Diabetes (1.4)	Skin (0.87)	Migraine (0.97)	Iron (1.11)	Oral (1.06)	Anxiety (0.81)	Disaster (138.88)
Azerbaijan	Back & neck (0.98)	Sense (1.08)	Depression (1.02)	Iron (1.55)	Iron (0.88)	Diabetes (1.36)	Migraine (0.96)	Anxiety (0.82)	Oral (1.03)	Osteoarthritis (1.26)
Georgia	Back & neck (0.93)	Sense (1.05)	Depression (1.01)	Diabetes (1.19)	Skin (0.85)	Migraine (0.98)	Iron (1.25)	Oral (1.02)	Anxiety (0.82)	Osteoarthritis (1.21)

(Figure 7 continues on next page)

	1	2	3	4	5	6	7	8	9	10
Kazakhstan	Back & neck (1-04)	Iron (2-4)	Sense (1-1)	Depression (1-03)	Skin (0-9)	Migraine (0-96)	Diabetes (1-17)	Oral (1-14)	Anxiety (0-82)	Osteoarthritis (1-31)
Kyrgyzstan	Back & neck (1-04)	Iron (1-3)	Depression (0-91)	Sense (0-85)	Skin (0-86)	Migraine (1-0)	Anxiety (0-76)	Diabetes (0-52)	Oral (1-05)	Asthma (0-69)
Mongolia	Back & neck (0-98)	Depression (0-94)	Sense (0-94)	Skin (0-87)	Iron (0-97)	Migraine (0-98)	Diabetes (0-79)	Anxiety (0-78)	Oral (1-02)	Falls (1-37)
Tajikistan	Back & neck (1-06)	Iron (1-03)	Depression (0-87)	Sense (0-8)	Skin (0-85)	Migraine (1-0)	Anxiety (0-78)	Diabetes (0-66)	Oral (1-06)	Epilepsy (1-54)
Turkmenistan	Back & neck (0-94)	Depression (1-0)	Sense (1-06)	Iron (1-31)	Skin (0-88)	Migraine (0-96)	Diabetes (1-33)	Anxiety (0-82)	Oral (1-02)	Asthma (0-65)
Uzbekistan	Back & neck (1-0)	Depression (0-94)	Sense (0-93)	Iron (1-08)	Skin (0-87)	Migraine (0-98)	Diabetes (0-84)	Anxiety (0-78)	Oral (1-03)	Asthma (0-79)
Latin America and Caribbean	Back & neck (0-87)	Sense (1-06)	Depression (1-08)	Skin (0-99)	Anxiety (1-52)	Iron (1-02)	Diabetes (1-12)	Migraine (1-0)	Asthma (1-18)	Other MSK (1-01)
Central Latin America	Back & neck (0-86)	Sense (1-12)	Depression (0-94)	Diabetes (1-44)	Skin (0-95)	Iron (0-93)	Migraine (0-9)	Anxiety (1-01)	Other MSK (0-89)	Oral (1-08)
Colombia	Back & neck (0-94)	Sense (1-14)	Depression (1-02)	Skin (1-02)	Anxiety (1-35)	Migraine (0-89)	Diabetes (0-81)	Asthma (1-23)	Other MSK (0-93)	Oral (1-07)
Costa Rica	Back & neck (0-89)	Sense (1-16)	Depression (1-01)	Skin (0-91)	Iron (1-01)	Migraine (0-88)	Anxiety (1-08)	Diabetes (0-72)	Asthma (1-18)	Other MSK (0-98)
El Salvador	Back & neck (0-94)	Sense (0-96)	Iron (1-59)	Depression (0-9)	Skin (0-94)	Diabetes (0-94)	Migraine (0-91)	Anxiety (1-03)	Asthma (1-24)	Oral (1-07)
Guatemala	Back & neck (0-96)	Iron (1-06)	Sense (0-94)	Skin (0-96)	Depression (0-85)	Diabetes (1-67)	Migraine (0-9)	Anxiety (1-05)	Asthma (1-02)	Oral (1-18)
Honduras	Back & neck (0-98)	Sense (0-95)	Skin (1-06)	Depression (0-88)	Iron (0-79)	Asthma (1-54)	Migraine (0-91)	Anxiety (1-05)	Diabetes (0-89)	Oral (1-07)
Mexico	Back & neck (0-79)	Sense (1-17)	Diabetes (1-9)	Depression (0-91)	Skin (0-91)	Migraine (0-91)	Anxiety (0-85)	Other MSK (0-94)	Oral (1-08)	Iron (0-5)
Nicaragua	Back & neck (0-96)	Sense (0-95)	Depression (0-87)	Skin (0-96)	Iron (0-82)	Migraine (0-91)	Anxiety (1-05)	Diabetes (0-85)	Asthma (0-96)	Oral (1-08)
Panama	Back & neck (0-91)	Sense (1-19)	Depression (0-99)	Skin (0-95)	Diabetes (1-35)	Iron (1-2)	Migraine (0-88)	Asthma (1-37)	Anxiety (1-09)	Oral (1-03)
Venezuela	Iron (3-06)	Back & neck (0-9)	Sense (1-12)	Depression (0-96)	Skin (0-98)	Diabetes (1-17)	Migraine (0-86)	Anxiety (1-06)	Asthma (1-01)	Other MSK (0-9)
Andean Latin America	Back & neck (0-93)	Sense (1-07)	Depression (1-09)	Iron (1-34)	Skin (1-02)	Anxiety (1-37)	Migraine (1-0)	Diabetes (0-76)	Asthma (1-1)	Oral (1-09)
Bolivia	Back & neck (1-0)	Iron (1-52)	Sense (0-97)	Depression (1-03)	Skin (1-02)	Anxiety (1-33)	Migraine (0-99)	Diabetes (0-73)	Asthma (1-11)	Oral (1-12)
Ecuador	Back & neck (0-91)	Sense (1-02)	Depression (1-08)	Iron (1-33)	Skin (0-99)	Diabetes (1-15)	Anxiety (1-37)	Migraine (0-87)	Oral (1-09)	Asthma (0-86)
Peru	Back & neck (0-92)	Sense (1-14)	Depression (1-11)	Skin (1-03)	Iron (1-29)	Migraine (1-07)	Anxiety (1-38)	Asthma (1-22)	Oral (1-09)	Diabetes (0-57)
Caribbean	Back & neck (0-89)	Sense (1-06)	Depression (1-11)	Iron (1-2)	Skin (1-09)	Diabetes (1-24)	Anxiety (1-49)	Migraine (1-0)	Asthma (1-62)	Other MSK (1-05)
Antigua	Back & neck (0-82)	Depression (1-2)	Sense (1-21)	Diabetes (2-2)	Skin (1-05)	Iron (1-43)	Anxiety (1-49)	Migraine (0-94)	Other MSK (1-37)	Asthma (1-31)
Bahamas	Back & neck (0-82)	Sense (1-19)	Depression (1-18)	Diabetes (1-94)	Skin (1-05)	Iron (1-54)	Anxiety (1-49)	Migraine (0-94)	Other MSK (1-47)	Asthma (1-38)
Barbados	Back & neck (0-81)	Sense (1-07)	Diabetes (2-05)	Depression (1-15)	Skin (0-96)	Other MSK (1-57)	Iron (1-49)	Anxiety (1-46)	Migraine (0-94)	Asthma (1-26)
Belize	Back & neck (0-84)	Depression (1-04)	Iron (1-19)	Skin (1-05)	Sense (0-92)	Anxiety (1-35)	Diabetes (1-14)	Migraine (0-97)	Asthma (1-58)	Other MSK (1-13)
Bermuda	Back & neck (0-89)	Depression (1-25)	Sense (1-19)	Skin (1-02)	Iron (1-32)	Anxiety (1-53)	Diabetes (1-87)	Migraine (0-96)	Other MSK (1-41)	Asthma (1-46)
Cuba	Back & neck (0-82)	Sense (1-11)	Depression (1-14)	Skin (1-08)	Diabetes (0-91)	Anxiety (1-46)	Iron (1-36)	Migraine (0-95)	Other MSK (1-11)	Asthma (1-31)
Dominica	Back & neck (0-83)	Diabetes (2-12)	Sense (1-07)	Depression (1-12)	Skin (1-05)	Iron (1-29)	Anxiety (1-41)	Migraine (0-95)	Asthma (1-52)	Other MSK (1-25)
Dominican Republic	Back & neck (0-82)	Sense (0-98)	Depression (1-07)	Skin (1-05)	Iron (1-19)	Anxiety (1-37)	Migraine (0-97)	Asthma (1-5)	Diabetes (0-71)	Other MSK (0-91)
Grenada	Back & neck (0-8)	Depression (1-11)	Sense (1-06)	Iron (1-52)	Skin (1-05)	Diabetes (1-78)	Other MSK (1-73)	Anxiety (1-41)	Migraine (0-94)	Asthma (1-43)
Guyana	Back & neck (0-84)	Iron (1-41)	Depression (1-02)	Sense (0-91)	Diabetes (1-36)	Skin (1-04)	Anxiety (1-33)	Migraine (0-97)	Asthma (1-58)	Other MSK (0-85)
Haiti	Iron (1-01)	Back & neck (0-96)	Depression (0-89)	Disaster (1227-89)	Skin (1-02)	Sense (0-8)	Asthma (1-97)	Anxiety (1-45)	Migraine (1-0)	Diabetes (1-38)
Jamaica	Back & neck (0-83)	Sense (1-0)	Depression (1-1)	Diabetes (1-43)	Skin (1-05)	Iron (1-25)	Anxiety (1-38)	Migraine (0-95)	Other MSK (1-39)	Asthma (1-48)
Puerto Rico	Back & neck (0-88)	Sense (1-2)	Diabetes (2-37)	Depression (1-23)	Skin (1-02)	Anxiety (1-53)	Iron (1-38)	Other MSK (1-3)	Migraine (0-95)	Asthma (1-41)
Saint Lucia	Back & neck (0-83)	Sense (1-05)	Depression (1-11)	Diabetes (1-48)	Skin (1-05)	Iron (1-26)	Anxiety (1-41)	Migraine (0-95)	Asthma (1-52)	Other MSK (1-25)
Saint Vincent and the Grenadines	Back & neck (0-82)	Sense (1-05)	Depression (1-1)	Diabetes (1-84)	Skin (1-05)	Iron (1-23)	Anxiety (1-41)	Migraine (0-95)	Asthma (1-45)	Other MSK (1-21)

(Figure 7 continues on next page)

	1	2	3	4	5	6	7	8	9	10
Suriname	Back & neck (0.82)	Sense (0.98)	Depression (1.07)	Skin (1.05)	Iron (1.25)	Diabetes (1.21)	Anxiety (1.38)	Migraine (0.96)	Asthma (1.53)	Other MSK (1.19)
Trinidad and Tobago	Back & neck (0.81)	Diabetes (3.11)	Sense (1.17)	Depression (1.18)	Skin (1.11)	Iron (1.44)	Anxiety (1.5)	Other MSK (1.49)	Migraine (0.94)	Asthma (1.58)
Virgin Islands, USA	Back & neck (0.89)	Sense (1.19)	Diabetes (2.14)	Depression (1.24)	Skin (1.02)	Other MSK (1.44)	Iron (1.38)	Anxiety (1.54)	Migraine (0.96)	Asthma (1.46)
Tropical Latin America	Back & neck (0.87)	Depression (1.23)	Sense (0.99)	Anxiety (2.14)	Skin (1.0)	Migraine (1.12)	Iron (1.0)	Diabetes (0.86)	Other MSK (1.21)	Asthma (1.41)
Brazil	Back & neck (0.86)	Depression (1.23)	Sense (0.99)	Anxiety (2.15)	Skin (1.0)	Migraine (1.12)	Iron (1.0)	Diabetes (0.86)	Other MSK (1.22)	Asthma (1.43)
Paraguay	Back & neck (0.97)	Depression (1.24)	Sense (0.97)	Skin (1.14)	Anxiety (1.81)	Iron (1.06)	Migraine (1.06)	Diabetes (0.92)	Other MSK (0.9)	Asthma (1.01)
Southeast Asia, east Asia, and Oceania	Back & neck (0.86)	Sense (0.94)	Depression (0.74)	Skin (1.0)	Diabetes (0.76)	Other MSK (1.22)	Iron (0.8)	Migraine (0.65)	Schiz (1.31)	Anxiety (0.72)
East Asia	Back & neck (0.89)	Sense (0.91)	Depression (0.73)	Skin (0.96)	Diabetes (0.64)	Other MSK (1.1)	Iron (0.79)	Schiz (1.38)	Migraine (0.57)	Anxiety (0.71)
China	Back & neck (0.88)	Sense (0.91)	Depression (0.74)	Skin (0.96)	Diabetes (0.62)	Other MSK (1.09)	Iron (0.79)	Schiz (1.39)	Migraine (0.57)	Anxiety (0.7)
North Korea	Back & neck (1.05)	Sense (0.77)	Skin (0.93)	Iron (0.92)	Depression (0.61)	Other MSK (1.15)	Diabetes (0.59)	COPD (2.1)	Anxiety (0.83)	Migraine (0.59)
Taiwan (province of China)	Back & neck (0.99)	Sense (1.08)	Diabetes (2.29)	Other MSK (1.59)	Depression (0.73)	Skin (0.89)	Anxiety (1.03)	Iron (1.05)	Migraine (0.63)	Osteoarthritis (1.44)
Southeast Asia	Back & neck (0.81)	Sense (1.01)	Skin (1.06)	Depression (0.76)	Diabetes (1.1)	Other MSK (1.53)	Iron (0.81)	Migraine (0.83)	Anxiety (0.75)	Asthma (1.02)
Cambodia	Back & neck (0.94)	Iron (0.94)	Sense (0.95)	Skin (1.05)	Depression (0.67)	Migraine (0.82)	Other MSK (1.11)	Asthma (1.14)	Anxiety (0.81)	Diabetes (0.82)
Indonesia	Back & neck (0.76)	Sense (0.99)	Skin (1.05)	Diabetes (1.17)	Depression (0.77)	Other MSK (1.67)	Iron (0.94)	Migraine (0.83)	Anxiety (0.77)	Asthma (0.93)
Laos	Back & neck (0.9)	Skin (1.05)	Iron (0.7)	Sense (0.84)	Depression (0.68)	Diabetes (1.36)	Other MSK (1.27)	Migraine (0.82)	Asthma (1.28)	Anxiety (0.8)
Malaysia	Back & neck (0.75)	Sense (1.0)	Skin (1.05)	Depression (0.84)	Diabetes (1.51)	Other MSK (1.82)	Nematode (4405.93)	Anxiety (1.2)	Migraine (0.79)	Schiz (1.18)
Maldives	Back & neck (0.82)	Sense (0.98)	Iron (1.35)	Skin (1.08)	Depression (0.75)	Migraine (0.84)	Other MSK (1.17)	Anxiety (0.76)	Haemog (3.61)	Diabetes (0.56)
Mauritius	Back & neck (0.82)	Diabetes (2.04)	Sense (1.15)	Depression (0.84)	Skin (1.07)	Other MSK (1.77)	Migraine (0.82)	Anxiety (0.8)	Asthma (1.03)	Oral (0.84)
Myanmar	Sense (1.13)	Back & neck (0.79)	Skin (1.04)	Depression (0.69)	Diabetes (1.23)	Iron (0.69)	Other MSK (1.38)	Migraine (0.82)	Asthma (1.08)	Anxiety (0.79)
Philippines	Back & neck (0.95)	Sense (1.05)	Skin (1.07)	Diabetes (1.49)	Iron (0.92)	Depression (0.75)	Other MSK (1.69)	Migraine (0.82)	Asthma (1.29)	Anxiety (0.77)
Sri Lanka	Back & neck (0.81)	Sense (1.14)	Diabetes (1.44)	Skin (1.06)	Depression (0.81)	Iron (1.04)	Migraine (0.82)	Asthma (1.14)	Anxiety (0.79)	Other MSK (0.86)
Seychelles	Back & neck (0.79)	Sense (1.18)	Skin (1.07)	Depression (0.84)	Diabetes (1.34)	Migraine (0.82)	Other MSK (1.15)	Anxiety (0.81)	Asthma (0.91)	Schiz (1.15)
Thailand	Back & neck (0.73)	Sense (1.04)	Depression (0.82)	Skin (1.06)	Other MSK (1.63)	Diabetes (0.83)	Migraine (0.84)	Iron (0.82)	Anxiety (0.79)	Asthma (1.08)
Timor-Leste	Back & neck (0.87)	Sense (0.97)	Iron (0.66)	Skin (1.07)	Depression (0.7)	Migraine (0.82)	Asthma (1.15)	Other MSK (1.03)	Diabetes (1.05)	Anxiety (0.84)
Vietnam	Back & neck (0.85)	Sense (0.92)	Skin (1.07)	Depression (0.77)	Migraine (0.85)	Other MSK (1.18)	Diabetes (0.6)	Schiz (1.2)	Other nutr (72.36)	Asthma (0.86)
Oceania	Back & neck (1.01)	Iron (0.98)	Skin (1.36)	Diabetes (2.69)	Sense (0.96)	Depression (0.7)	Asthma (1.76)	Other MSK (1.58)	Nematode (3.75)	Migraine (0.77)
American Samoa	Back & neck (0.9)	Diabetes (2.78)	Skin (1.34)	Sense (1.03)	Iron (0.91)	Depression (0.77)	Other MSK (1.87)	Asthma (1.31)	Migraine (0.72)	Anxiety (0.83)
Micronesia	Back & neck (0.91)	Skin (1.31)	Diabetes (1.68)	Sense (0.91)	Depression (0.72)	Other MSK (1.65)	Asthma (1.45)	Iron (0.66)	Migraine (0.73)	Anxiety (0.82)
Fiji	Diabetes (3.07)	Back & neck (0.84)	Skin (1.27)	Sense (1.03)	Iron (1.1)	Depression (0.75)	Asthma (1.64)	Other MSK (1.45)	Nematode (58.08)	Migraine (0.72)
Guam	Back & neck (0.94)	Diabetes (3.17)	Sense (1.27)	Skin (1.24)	Other MSK (1.96)	Depression (0.89)	Iron (0.97)	Asthma (1.47)	Other NTD (234.964.29)	Migraine (0.72)
Kiribati	Back & neck (0.93)	Diabetes (3.07)	Nematode (7.02)	Skin (1.27)	Sense (0.81)	Iron (0.54)	Depression (0.65)	Asthma (1.68)	Migraine (0.72)	Other MSK (1.03)
Marshall	Diabetes (3.31)	Back & neck (0.88)	Skin (1.29)	Iron (1.23)	Sense (0.87)	Nematode (7.21)	Depression (0.69)	Other MSK (1.7)	Asthma (1.41)	Migraine (0.72)
Northern Mariana Islands	Back & neck (0.86)	Skin (1.25)	Diabetes (3.3)	Depression (0.82)	Sense (1.28)	Other MSK (1.61)	Iron (1.08)	Migraine (0.71)	Anxiety (0.85)	Asthma (1.16)
Papua New Guinea	Iron (0.97)	Back & neck (0.95)	Skin (1.29)	Sense (0.84)	Diabetes (2.35)	Depression (0.63)	Asthma (1.7)	Nematode (3.34)	Other MSK (1.43)	LF (100.35)
Samoa	Back & neck (0.9)	Skin (1.33)	Sense (0.92)	Diabetes (1.5)	Iron (0.84)	Depression (0.74)	Other MSK (1.6)	Asthma (1.16)	Migraine (0.73)	Anxiety (0.82)
Solomon Islands	Back & neck (0.95)	Skin (1.25)	Diabetes (2.69)	Sense (0.81)	Iron (0.53)	Depression (0.65)	Other MSK (1.56)	Asthma (1.45)	Migraine (0.73)	Anxiety (0.88)
Tonga	Back & neck (0.92)	Skin (1.34)	Diabetes (1.49)	Sense (0.83)	Iron (0.97)	Depression (0.73)	Other MSK (1.69)	Asthma (1.68)	Migraine (0.73)	Osteoarthritis (2.01)

(Figure 7 continues on next page)

	1	2	3	4	5	6	7	8	9	10
Vanuatu	Back & neck (0-91)	Skin (1-31)	Diabetes (1-79)	Sense (0-82)	Depression (0-67)	Other MSK (1-55)	Asthma (1-45)	Migraine (0-73)	Nematode (2-91)	Anxiety (0-83)
North Africa and Middle East	Back & neck (1-07)	Depression (0-93)	Sense (0-86)	Diabetes (1-62)	Iron (0-82)	Skin (0-84)	Migraine (1-07)	Other MSK (1-34)	Anxiety (1-06)	Asthma (0-94)
North Africa and Middle East	Back & neck (1-07)	Depression (0-93)	Sense (0-86)	Diabetes (1-62)	Iron (0-82)	Skin (0-84)	Migraine (1-07)	Other MSK (1-34)	Anxiety (1-06)	Asthma (0-94)
Afghanistan	Iron (0-8)	Back & neck (1-15)	War (163-45)	Skin (0-84)	Depression (0-8)	Sense (0-73)	Diabetes (2-72)	Other MSK (5-28)	Migraine (1-18)	Anxiety (1-23)
Algeria	Back & neck (1-04)	Diabetes (1-47)	Depression (0-95)	Sense (0-82)	Skin (0-83)	Migraine (1-08)	Other MSK (1-52)	Iron (0-7)	Anxiety (1-04)	Asthma (1-0)
Bahrain	Back & neck (0-95)	Diabetes (2-9)	Depression (1-11)	Sense (1-02)	Other MSK (1-96)	Skin (0-9)	Migraine (1-07)	Iron (1-43)	Anxiety (1-07)	Drugs (1-78)
Egypt	Back & neck (1-08)	Iron (1-53)	Diabetes (1-5)	Sense (0-93)	Depression (0-77)	Skin (0-84)	Migraine (1-07)	Other MSK (1-19)	Anxiety (1-0)	Asthma (0-96)
Iran	Back & neck (1-03)	Depression (1-05)	Sense (0-94)	Diabetes (1-48)	Migraine (0-98)	Skin (0-8)	Other MSK (1-44)	Anxiety (1-07)	Drugs (3-41)	Other cardio (6-21)
Iraq	Back & neck (1-08)	Diabetes (2-19)	Depression (0-93)	War (395-43)	Sense (0-79)	Skin (0-75)	Iron (0-65)	Migraine (1-06)	Anxiety (1-14)	Other MSK (0-92)
Jordan	Back & neck (1-0)	Depression (1-01)	Diabetes (1-99)	Skin (0-92)	Iron (0-97)	Sense (0-92)	Migraine (1-05)	Other MSK (1-69)	Anxiety (1-07)	Asthma (1-03)
Kuwait	Back & neck (0-88)	Depression (1-18)	Migraine (1-06)	Skin (0-89)	Diabetes (2-46)	Sense (1-09)	Iron (1-18)	Anxiety (1-12)	Other MSK (1-32)	Asthma (1-01)
Lebanon	War (10689-82)	Back & neck (0-83)	Diabetes (2-0)	Depression (1-02)	Sense (0-9)	Skin (0-93)	Iron (1-35)	Migraine (1-03)	Anxiety (1-27)	Asthma (1-14)
Libya	Back & neck (1-04)	Depression (1-0)	Diabetes (1-3)	Sense (0-81)	Skin (0-87)	Migraine (1-07)	War (464-77)	Iron (0-72)	Anxiety (1-04)	Other MSK (1-19)
Morocco	Back & neck (1-14)	Diabetes (2-21)	Sense (0-79)	Depression (0-88)	Skin (0-92)	Migraine (1-05)	Iron (0-56)	Other MSK (1-36)	Anxiety (1-09)	Drugs (4-01)
Palestine	Back & neck (1-06)	Depression (1-29)	Skin (0-83)	Sense (0-72)	Iron (0-59)	Migraine (1-07)	Anxiety (1-07)	Diabetes (1-14)	War (198-72)	Other MSK (1-08)
Oman	Back & neck (0-95)	Depression (1-07)	Diabetes (2-42)	Other Cardio (19-2)	Sense (1-01)	Migraine (1-1)	Skin (0-85)	Iron (1-15)	Other MSK (1-43)	Anxiety (1-05)
Qatar	Back & neck (0-94)	Depression (1-15)	Diabetes (3-12)	Migraine (1-12)	Skin (0-94)	Sense (0-96)	Anxiety (1-05)	Other MSK (1-26)	Iron (1-14)	Heat & cold (17-26)
Saudi Arabia	Back & neck (0-99)	Depression (1-11)	Migraine (1-27)	Skin (0-93)	Sense (1-01)	Diabetes (1-51)	Anxiety (1-08)	Other MSK (1-33)	Iron (0-52)	Asthma (0-8)
Sudan	Back & neck (1-07)	Iron (0-81)	Depression (0-84)	Skin (0-84)	Sense (0-75)	Migraine (1-08)	Diabetes (1-9)	Anxiety (1-15)	Asthma (1-18)	Other MSK (1-16)
Syria	War (1612-15)	Back & neck (1-05)	Depression (0-94)	Sense (0-78)	Skin (0-8)	Iron (0-73)	Migraine (1-07)	Diabetes (1-07)	Anxiety (1-06)	Asthma (0-99)
Tunisia	Back & neck (1-02)	Diabetes (1-35)	Depression (1-0)	Sense (0-82)	Skin (0-89)	Other MSK (1-58)	Migraine (1-02)	Anxiety (1-04)	Iron (0-66)	Asthma (1-05)
Turkey	Back & neck (1-13)	Sense (0-85)	Depression (0-92)	Diabetes (1-3)	Other MSK (1-67)	Skin (0-83)	Migraine (1-01)	Iron (0-93)	Anxiety (0-92)	Oral (1-18)
United Arab Emirates	Back & neck (0-99)	Depression (1-18)	Diabetes (3-52)	Other MSK (1-92)	Migraine (1-14)	Sense (1-15)	Skin (0-91)	Drugs (2-16)	Anxiety (1-04)	Iron (1-21)
Yemen	Back & neck (1-11)	War (248-22)	Depression (0-84)	Skin (0-84)	Sense (0-77)	Iron (0-44)	Migraine (1-11)	Diabetes (1-76)	Anxiety (1-17)	Other MSK (1-34)
South Asia	Iron (1-37)	Back & neck (0-91)	Sense (1-11)	Depression (0-92)	Other MSK (1-96)	Skin (0-89)	Migraine (1-19)	Diabetes (0-9)	Anxiety (0-8)	COPD (1-77)
South Asia	Iron (1-37)	Back & neck (0-91)	Sense (1-11)	Depression (0-92)	Other MSK (1-96)	Skin (0-89)	Migraine (1-19)	Diabetes (0-9)	Anxiety (0-8)	COPD (1-77)
Bangladesh	Back & neck (1-06)	Other MSK (2-7)	Sense (0-95)	Depression (0-82)	Iron (0-67)	Skin (0-94)	Migraine (1-2)	Anxiety (1-11)	Diabetes (1-0)	Epilepsy (2-97)
Bhutan	Back & neck (1-05)	Iron (1-48)	Sense (0-98)	Depression (0-83)	Skin (0-95)	Other MSK (1-84)	Migraine (1-19)	Diabetes (1-17)	Diarrhoea (3-33)	Anxiety (0-88)
India	Iron (1-58)	Back & neck (0-89)	Sense (1-16)	Depression (0-94)	Other MSK (1-93)	Migraine (1-2)	Skin (0-86)	Diabetes (0-83)	Anxiety (0-74)	COPD (2-01)
Nepal	Back & neck (1-13)	Sense (0-89)	Skin (0-97)	Migraine (1-33)	Other MSK (1-93)	Iron (0-48)	Depression (0-63)	Disaster (661-96)	Anxiety (0-94)	Asthma (1-07)
Pakistan	Iron (1-02)	Back & neck (0-87)	Sense (0-93)	Depression (0-9)	Skin (1-0)	Migraine (1-13)	Other MSK (1-56)	Diabetes (1-47)	Anxiety (0-92)	Asthma (0-91)
Sub-Saharan Africa	Iron (0-79)	Back & neck (0-98)	Depression (1-04)	Sense (1-02)	Skin (0-93)	Other NTD (1-24)	Migraine (0-8)	HIV (4-74)	Asthma (0-98)	Malaria (2-25)
Southern sub-Saharan Africa	HIV (111-59)	Back & neck (1-03)	Depression (1-05)	Sense (1-06)	Skin (0-95)	Iron (0-96)	Diabetes (1-54)	Migraine (0-78)	Asthma (1-29)	Anxiety (0-79)
Botswana	HIV (140-61)	Back & neck (0-98)	Iron (1-51)	Depression (1-09)	Sense (0-98)	Skin (0-95)	Diabetes (1-33)	Asthma (1-49)	Migraine (0-77)	TB (15-89)
Lesotho	HIV (73-48)	Back & neck (1-08)	Depression (1-14)	Sense (0-91)	Skin (0-93)	Iron (0-64)	Diabetes (1-63)	Asthma (1-68)	Migraine (0-77)	Anxiety (0-78)
Namibia	HIV (77-97)	Back & neck (0-99)	Iron (1-28)	Depression (1-09)	Sense (0-95)	Skin (0-96)	Diabetes (1-07)	Asthma (1-42)	Migraine (0-78)	Anxiety (0-76)
South Africa	HIV (165-51)	Back & neck (1-04)	Sense (1-12)	Depression (1-04)	Diabetes (1-65)	Skin (0-95)	Iron (1-01)	Migraine (0-78)	Asthma (1-27)	Anxiety (0-81)

(Figure 7 continues on next page)

	1	2	3	4	5	6	7	8	9	10
Swaziland	HIV (172-35)	Back & neck (0-94)	Depression (1-09)	Skin (0-95)	Sense (0-97)	Iron (0-92)	Diabetes (1-51)	Asthma (1-95)	Nematode (5-73)	Migraine (0-77)
Zimbabwe	HIV (46-19)	Back & neck (1-04)	Iron (0-82)	Depression (1-04)	Skin (0-94)	Sense (0-88)	Asthma (1-18)	Migraine (0-78)	Diabetes (1-12)	Anxiety (0-75)
Western sub-Saharan Africa	Iron (0-99)	Back & neck (1-11)	Depression (0-99)	Sense (0-98)	Skin (0-85)	Malaria (3-39)	Migraine (0-79)	Schisto (10-95)	Anxiety (0-82)	Asthma (0-82)
Benin	Iron (0-76)	Back & neck (1-08)	Depression (0-94)	Sense (0-91)	Skin (0-91)	Malaria (5-86)	Migraine (0-85)	Anxiety (0-83)	Asthma (0-81)	Schisto (43-81)
Burkina Faso	Iron (0-73)	Back & neck (1-08)	Depression (0-91)	Skin (0-92)	Sense (0-87)	Malaria (1-56)	Migraine (0-9)	Haemog (45-65)	Anxiety (0-9)	Diarrhoea (0-81)
Cameroon	Back & neck (1-05)	Depression (0-99)	Iron (0-57)	Skin (0-92)	Sense (0-93)	Oncho (654332-73)	HIV (9-44)	Malaria (45-65)	Migraine (0-77)	Anxiety (0-78)
Cape Verde	Back & neck (1-01)	Depression (1-05)	Iron (1-08)	Sense (0-98)	Skin (0-93)	Migraine (0-76)	Diabetes (0-83)	Other cardio (9-52)	Anxiety (0-73)	Asthma (0-76)
Chad	Iron (1-31)	Back & neck (1-18)	Skin (0-91)	Depression (0-92)	Sense (0-92)	Migraine (0-86)	Diarrhoea (1-0)	Asthma (0-85)	Anxiety (0-87)	Heat & cold (4-25)
Côte d'Ivoire	Iron (0-75)	Back & neck (1-06)	Skin (1-08)	Sense (0-97)	Depression (0-94)	Malaria (15-32)	Migraine (0-8)	HIV (4-5)	Asthma (0-91)	Anxiety (0-81)
The Gambia	Iron (0-73)	Back & neck (1-08)	Depression (1-01)	Skin (0-91)	Sense (0-84)	Malaria (3-43)	Migraine (0-84)	Anxiety (0-85)	Diarrhoea (0-94)	Asthma (0-74)
Ghana	Back & neck (0-92)	Iron (0-87)	Depression (1-0)	Sense (0-99)	Skin (0-69)	Malaria (150-78)	Migraine (0-76)	Schisto (6627-12)	Anxiety (0-75)	Diabetes (0-66)
Guinea	Back & neck (1-1)	Iron (0-68)	Skin (1-06)	Sense (0-93)	Depression (0-92)	Malaria (2-14)	Schisto (13-79)	Migraine (0-87)	Asthma (0-95)	Anxiety (0-86)
Guinea-Bissau	Iron (0-77)	Back & neck (1-11)	Depression (0-92)	Sense (0-92)	Skin (0-91)	Migraine (0-85)	HIV (1-77)	Schisto (2-78)	Schisto (15-56)	Asthma (0-9)
Liberia	Oncho (92161-85)	Back & neck (1-1)	Iron (0-73)	Sense (0-91)	Skin (0-91)	Depression (0-82)	Schisto (18-67)	Malaria (1-85)	Migraine (0-85)	Anxiety (0-85)
Mali	Iron (1-11)	Back & neck (0-93)	Sense (0-94)	Depression (0-92)	Skin (0-83)	Malaria (1-11)	Migraine (0-9)	Anxiety (0-9)	Heat & cold (3-29)	Diarrhoea (0-66)
Mauritania	Iron (1-24)	Back & neck (1-07)	Sense (1-06)	Depression (0-96)	Skin (0-91)	Migraine (0-79)	Asthma (0-94)	Anxiety (0-8)	Heat & cold (5-99)	Schisto (237-73)
Niger	Iron (0-78)	Back & neck (1-16)	Sense (0-94)	Skin (0-94)	Depression (0-92)	Migraine (0-98)	Heat & cold (3-9)	Malaria (0-32)	Diarrhoea (0-77)	Anxiety (0-97)
Nigeria	Iron (1-18)	Back & neck (1-18)	Depression (1-05)	Sense (1-03)	Skin (0-78)	Schisto (2926-68)	Malaria (58-71)	Migraine (0-73)	Anxiety (0-8)	Asthma (0-86)
São Tomé and Príncipe	Back & neck (1-04)	Iron (0-65)	Depression (0-99)	Sense (0-97)	Skin (0-92)	Other NTD (2-54)	Migraine (0-78)	Malaria (24-3)	Asthma (0-88)	Anxiety (0-79)
Senegal	Iron (1-04)	Back & neck (0-98)	Depression (0-95)	Sense (0-93)	Skin (0-91)	Migraine (0-83)	Anxiety (0-84)	Diarrhoea (0-96)	Asthma (0-72)	Diabetes (1-01)
Sierra Leone	Back & neck (1-08)	Iron (0-59)	Depression (0-93)	Skin (0-91)	Sense (0-87)	Malaria (5-16)	Migraine (0-84)	Oncho (46362-69)	Asthma (0-86)	Anxiety (0-84)
Togo	Back & neck (1-1)	Iron (0-71)	Depression (0-95)	Sense (0-97)	Skin (0-94)	Malaria (8-85)	Migraine (0-82)	Asthma (0-93)	Anxiety (0-82)	Diarrhoea (1-04)
Eastern sub-Saharan Africa	Depression (1-11)	Iron (0-62)	Sense (0-99)	Skin (0-95)	Back & neck (0-78)	Migraine (0-79)	Other NTD (0-88)	Anxiety (0-94)	HIV (3-82)	Asthma (0-94)
Burundi	Depression (1-05)	Back & neck (0-85)	Skin (0-92)	Sense (0-9)	Iron (0-26)	Migraine (0-88)	Asthma (1-0)	Diarrhoea (0-99)	Anxiety (1-0)	Malaria (0-79)
Comoros	Sense (1-09)	Depression (1-06)	Back & neck (0-86)	Iron (0-6)	Skin (0-91)	Malaria (7-6)	Migraine (0-8)	Asthma (1-06)	Anxiety (0-92)	Diabetes (1-11)
Djibouti	Iron (0-88)	Depression (1-12)	Sense (1-02)	Back & neck (0-73)	Skin (0-91)	Other NTD (2-76)	Heat & cold (9-43)	Migraine (0-75)	Diabetes (1-16)	Anxiety (0-87)
Eritrea	Iron (0-89)	Depression (1-06)	Sense (1-0)	Skin (0-91)	Back & neck (0-7)	War (102-99)	Schisto (42-57)	Migraine (0-81)	Asthma (0-97)	Anxiety (0-94)
Ethiopia	Depression (1-13)	Sense (1-0)	Skin (1-02)	Back & neck (0-8)	Iron (0-32)	Schisto (26-64)	Anxiety (1-0)	Migraine (0-76)	Other NTD (0-66)	Asthma (0-79)
Kenya	Iron (1-05)	Other NTD (4-38)	Depression (1-13)	Sense (1-09)	Skin (1-01)	Back & neck (0-81)	HIV (9-68)	Migraine (0-8)	Anxiety (0-87)	Schisto (17-62)
Madagascar	Iron (0-7)	Sense (1-14)	Depression (1-09)	Back & neck (0-83)	Skin (0-9)	Asthma (1-28)	Schisto (145-77)	Migraine (0-79)	Anxiety (0-92)	Diarrhoea (1-07)
Malawi	Depression (1-05)	Iron (0-53)	Back & neck (0-85)	Sense (0-93)	Skin (0-91)	HIV (7-57)	Malaria (2-62)	Migraine (0-83)	Anxiety (0-96)	Asthma (0-9)
Mozambique	Iron (0-71)	Depression (1-04)	Sense (0-99)	HIV (7-43)	Back & neck (0-83)	Skin (0-89)	Malaria (1-88)	Migraine (0-84)	Asthma (0-96)	Anxiety (0-97)
Rwanda	Back & neck (0-8)	Sense (0-94)	Depression (0-88)	Iron (0-5)	War (198-09)	Skin (0-77)	Asthma (1-25)	Migraine (0-79)	Anxiety (0-92)	Other NTD (0-82)
Somalia	Iron (1-23)	Depression (1-03)	Sense (0-99)	Skin (0-93)	Back & neck (0-81)	Schisto (1-65)	Iodine (3-5)	Asthma (1-03)	Migraine (0-93)	Other NTD (0-84)
South Sudan	Iron (0-81)	Depression (1-05)	Sense (1-02)	Oncho (45415-75)	Back & neck (0-81)	Skin (0-91)	Schisto (11-22)	Heat & cold (5-89)	Asthma (1-1)	Migraine (0-84)
Tanzania	Iron (0-85)	Depression (1-08)	Skin (0-93)	Sense (0-86)	Back & neck (0-65)	Migraine (0-72)	Asthma (0-97)	Anxiety (0-9)	HIV (5-38)	Malaria (12-32)
Uganda	Depression (1-31)	Skin (0-91)	Sense (0-96)	Iron (0-45)	Back & neck (0-79)	Malaria (9-75)	HIV (6-86)	Asthma (1-01)	Migraine (0-79)	Anxiety (0-89)

(Figure 7 continues on next page)

	1	2	3	4	5	6	7	8	9	10
Zambia	Depression (1.11)	HIV (20.66)	Skin (0.91)	Sense (0.96)	Iron (0.52)	Back & neck (0.77)	Migraine (0.88)	Schisto (1984.48)	Asthma (0.93)	Anxiety (0.88)
Central sub-Saharan Africa	Other NTD (4.36)	Iron (0.7)	Back & neck (1.1)	Sense (1.22)	Skin (1.11)	Depression (0.97)	Oncho (31767.28)	Asthma (1.44)	Malaria (1.43)	Migraine (0.87)
Angola	Other NTD (4.05)	Skin (1.13)	Sense (1.25)	Back & neck (1.0)	Depression (1.0)	Iron (0.38)	Asthma (1.59)	Schisto (567.16)	Malaria (15.61)	Migraine (0.8)
Central African Republic	Iron (0.84)	Back & neck (1.12)	Sense (1.17)	Skin (1.1)	Depression (0.95)	Oncho (37527.86)	Asthma (1.46)	Other NTD (0.95)	HIV (3.4)	Migraine (0.87)
Congo (Brazzaville)	Iron (1.03)	Sense (1.27)	Back & neck (1.0)	Skin (1.16)	Other NTD (5.85)	Depression (1.04)	Asthma (1.25)	Diabetes (1.2)	Schisto (10909.69)	Migraine (0.77)
DR Congo	Other NTD (4.56)	Iron (0.76)	Back & neck (1.15)	Sense (1.21)	Skin (1.1)	Oncho (31562.22)	Depression (0.96)	Asthma (1.4)	Malaria (1.11)	Migraine (0.9)
Equatorial Guinea	Back & neck (0.95)	Sense (1.23)	Skin (1.14)	Depression (1.11)	Diabetes (1.22)	Iron (0.75)	Asthma (1.72)	Malaria (6683.41)	Migraine (0.78)	HIV (21.87)
Gabon	Other NTD (24.82)	Iron (1.58)	Sense (1.31)	Back & neck (0.94)	Depression (1.13)	Skin (1.1)	Schisto (55308.55)	Diabetes (1.14)	Asthma (1.62)	Migraine (0.77)

Colour key

0.0–0.81 0.81–0.88 0.88–0.94 0.94–0.99 0.99–1.05
 1.05–1.13 1.13–1.26 1.26–1.58 ≥1.58

Figure 7: Leading ten causes of years lived with disability (YLDs) with the ratio of observed years lived with disability (YLDs) to years lived with disability (YLDs) expected on the basis of SDI in 2015, by location

Shades of blue represent much lower observed YLDs than expected levels based on SDI, whereas red shows that observed YLDs exceed expected levels. Alzheimer's=Alzheimer's disease and other dementias. Back & neck=low back and neck pain. COPD=chronic obstructive pulmonary disease. Drugs=drug use disorders. Haemog=haemoglobinopathies and haemolytic anaemias. Heat & cold=environmental heat and cold exposure. IHD=ischaeamic heart disease. Iron=iron-deficiency anaemia. NTD=neglected tropical diseases. Oncho=onchocerciasis. Oral=oral disorders. Osteoarth=osteoarthritis. Other cardio=other cardiovascular and circulatory diseases. Other MSK=other musculoskeletal disorders. Other nutr=other nutritional deficiencies. Other unint=other unintentional injuries. PEM=protein-energy malnutrition. Schisto=schistosomiasis. Schiz=schizophrenia. Sense=sense organ diseases. Skin=skin and subcutaneous diseases. TB=tuberculosis.

malaria prevalence and incidence become available for all malaria-endemic countries outside of Africa.

Tuberculosis

We added new data from tuberculosis prevalence surveys done in Indonesia, Ghana, and several subnational locations in India. Our analysis of tuberculosis relied on prevalence surveys, case notifications and cause-specific mortality estimates. We used the expert judgment values for the case-detection rates (CDRs) from WHO as an initial guide of how much notifications need to be increased to reflect incidence of all tuberculosis, but relied on DisMod-MR 2.1 to find an estimate that is consistent with available prevalence and cause-specific mortality rates. We found that, particularly in older age groups, our estimated CDR often falls below the all-age CDR of WHO. In GBD 2013, we used a relative risk approach to predict the proportion of HIV-infected individuals with tuberculosis. In GBD 2015, we improved our modelling strategy by making use of more abundantly available data on the proportions of HIV infected cases among all tuberculosis cases from the WHO case notifications to separate out combined HIV and tuberculosis from all forms of tuberculosis. In our modelling of tuberculosis, we have not separately estimated the incidence and prevalence related to multidrug-resistant tuberculosis. Given the policy interest in multidrug-resistant tuberculosis, we plan to include estimates for multidrug-resistant tuberculosis in future rounds of the GBD. Our global tuberculosis (all-forms) incidence estimate (10.2 million [9.2–11.5 million] cases in 2015) is slightly higher than that of WHO (9.6 million cases in 2014), and we estimate a slightly larger fraction of combined HIV and tuberculosis (13.0%)

than WHO (12.0%). Our list of countries with high burden of tuberculosis is consistent with that of WHO, with a few exceptions. Afghanistan and Cambodia are lower in our estimates, and surpassed by Ukraine and Angola.

Cardiovascular diseases

Ageing and population growth have led to a growing number of people living with atherosclerotic vascular disease worldwide, despite the decrease in incident myocardial infarction and ischaemic stroke in high-income regions. Rising levels of obesity and diabetes in many countries makes this an area of major concern for global health. Increased attention will need to be directed toward this highly treatable set of disorders. Health systems will need to improve the delivery of cost-effective treatments such as blood pressure and cholesterol-lowering drugs while efforts continue to focus on decreasing tobacco smoking, improving diet, and increasing physical activity. Investments in universal primary care and public health campaigns need to be balanced against the need to improve access to emergency care, including the strengthening of pre-hospital systems and expanded access to revascularisation treatments.

Cancer

The International Agency for Research on Cancer last produced cancer estimates by country, age, sex, and cancer site for 2012 for the GLOBOCAN project.⁶⁰ The total estimated cancer incidence from GLOBOCAN for 2012 was 14.1 million individuals. By comparison, GBD 2015 estimated 18.6 million (18.0–19.4 million) new cases of cancer in 2015, which includes cases of non-melanoma skin cancer. GLOBOCAN used nine different

methods to estimate incidence, which precludes the calculation of uncertainty for those estimates. The GBD relies on the strength of the mortality estimates and transformation of the mortality estimates to incidence, taking into account uncertainty associated with both the mortality estimates and with the mortality to incidence ratios. Until high-quality cancer incidence is available in all countries, the GBD approach, which maximises the amount of data used to determine incidence estimates, is highly beneficial.

Vision loss

Globally, 34·3 million (30·7–38·0 million) people are blind, an additional 24·3 million (21·6–27·6 million) have severe vision impairment, 214 million (193–237 million) have moderate vision impairment, and 663 million (638–690 million) have near vision impairment in 2015. Combined, vision loss accounts for 24·5 million (17·0–34·5 million) YLDs and is the third largest impairment after anaemia (77·9 million [52·4–111·4 million] YLDs) and hearing loss (46·2 million [31·6–63·2 million] YLDs). YLDs from vision loss are slightly lower than those for anxiety disorders, which rank ninth worldwide. Largely due to ageing, substantial increases occurred from 2005 to 2015 in the number of people with blindness (increasing by 23·3% [21·9–24·6%]), severe visual impairment (24·3% [22·4–26·0%]), moderate visual impairment (22·0% [20·1–23·7%]), and near vision impairment (22·5% [20·9–24·0%]). Most causes of vision loss can be prevented or cured using cost-effective interventions.^{61,62}

Chronic kidney disease

Compared with our GBD 2013 estimates, we estimated global prevalence of chronic kidney disease to be lower by a third. Data for chronic kidney disease at younger ages are sparse. We increased the number of datapoints below the age of 30 years from 20 to 51, and that led to considerably lower prevalence estimates for these ages. We have used different strategies for estimating deaths versus YLDs secondary to chronic kidney disease due to other causes. In our cause of death analysis, deaths secondary to many of the causes typically categorised within the chronic kidney disease other category, such as cystic disease and other congenital renal diseases, were already counted under the primary disease following the principles of the ICD, and thus could not be recounted within the chronic kidney disease other category. In comparison, for the YLD analysis we counted chronic kidney disease from these causes in the chronic kidney disease other category. This difference in the classification of morbidity and mortality from chronic kidney disease other is driven by the underlying nature of the data on causes of death and prevalence. In future GBD versions, we plan to make explicit estimates of death and YLDs for polycystic kidney disease, an important component of the other category.

Emerging infectious diseases

Emerging and re-emerging infectious diseases pose a persistent yet dynamic challenge to global health⁶³ and to the GBD study. GBD 2015 has included Ebola virus disease in response to the devastating outbreak in west Africa starting in 2013,⁶⁴ whereas we have not captured the recent expansions of chikungunya⁶⁵ and Zika virus across the Americas,⁶⁶ for instance. Quantifying the effect of newly emerging epidemic diseases is difficult because the formal reporting of the disease events might considerably lag the first presentation of cases. As a result, contemporary events such as the substantial deviation from baseline levels in reported congenital abnormalities in Brazil in 2015,⁶⁷ now established to be associated with Zika virus,⁶⁸ will need to be incorporated into GBD 2016.

Advances in data and analysis

DisMod-MR

For GBD 2015, we improved the estimation for subnational units in DisMod-MR 2.1 compared to GBD 2013 and made substantive efforts to standardise the modelling approach across diseases. Three changes had an important effect on our estimates. First, the cascade was implemented such that national estimates were used to inform subnational estimation (in GBD 2013, each subnational unit was treated as a country within a GBD region). Second, wherever possible, we improved the linkage in the estimation between cause-specific mortality and disease incidence and prevalence by the systematic inclusion of estimated excess mortality data to provide more informative priors for each location in the geographical hierarchy. Third, in GBD 2010, and to a lesser degree in GBD 2013, many DisMod-MR models did not include fixed effects of country covariates for incidence or prevalence. Spatial heterogeneity was largely estimated from the random effects. For GBD 2015, we systematically tested the inclusion of more fixed effects in DisMod-MR 2.1 models including, where appropriate, variables related to the risk factor exposures we estimate for each disease.

Disability weights

For GBD 2015, we have not incorporated any new population-based data on disability weights; we have used the same disability weights as for GBD 2013. The present set of disability weights were based on population surveys in nine countries and an open internet survey. So far, these studies have not found systematic variation in disability weights across populations or within populations as a function of education, income per capita, or other variables. In future, we hope that more population surveys on disability weights will continue to be done to enrich the database for disability weight estimation. Given that we have not been able to collect disability weights in every location included in the GBD, it is possible that disability weights might vary systematically across communities. Even if this were to be confirmed empirically, for global standardised comparisons we would still use the global

average disability weights; however, country analyses might use local disability weights. Until there is evidence of systematic variation across communities, this remains a theoretical and not a practical consideration. Given the close correlation between the internet survey and the population-based surveys, we are considering the implementation of an open rolling internet survey to collect more data for disability weights including new or revised health-state descriptions.

GATHER compliance

Providing all documentation for data sources, establishing access to the datasets used in modelling, and posting the code has been a labour-intensive activity; GBD 2015 is compliant with the new GATHER guidelines as a result.¹⁷ Posting code, we believe, will stimulate other researchers to explore the methods used in GBD estimation of non-fatal outcomes and hopefully lead to suggestions for improved estimation. With a steadily growing set of co-investigators and a widening community interested in global health estimation, the enhanced transparency of GBD will improve the efficacy of the overall effort.

Using claims data

For the USA we used claims data for selected disorders. These data were particularly useful in the estimation of state-by-state prevalence for some disorders. Claims data, however, have important potential biases. Because of exclusion of some disadvantaged groups from health insurance, disease rates might be different for individuals that are covered compared with those from the general population.^{69,70} As the claims dataset included information from Medicaid and Medicare, the coverage schemes for low-income citizens and older adults (older than 65 years) in the USA, the bias toward underestimation might not be so great. Indeed, we found that for many diseases, such as asthma, diabetes, or rheumatoid arthritis, the claims data were consistent with high-quality survey estimates. A potential problem of overestimation exists because visits to rule out a diagnosis can be coded to the diagnosis. Others working with claims have counted diagnoses only if they had appeared at least twice during a year. Applying such a restriction would have led to rejection of 44–68% of skin disorders, such as acne, psoriasis, or scabies, where one would expect that even after a single visit a diagnosis can be reliably made. As we tend to adjust the claims data used in DisMod-MR 2.1 if a systematic bias is detected from population-based sources, such as the National Health and Nutrition Examination Survey or other quality surveys, we believe our estimates have not been influenced much by this problem. The potential for greater inclusion of claims data in the GBD analysis is large; in countries with universal health access, claims data might overcome the problems of non-responder bias in survey data, which tend to lead to underestimates of the true population prevalence or incidence.

Self-reported functional health status

We found only modest progress in high-SDI countries in reducing age-standardised YLDs per capita over the last 25 years. There are, however, reports in the literature of substantial improvements based on self-rated health.^{71,72} In the USA, these improvements seem to be survey programme-specific and have not been reported in all data systems.⁷¹ Nevertheless, more work is needed to understand the different trends between our assessment of YLDs per capita and self-rated health. Item response theory has been used in some fields to identify changing response patterns for single items compared with the pool of all items.⁷³ Item response theory cannot capture systematic changes that affect a set of items, and anchoring vignettes have been proposed as a strategy to deal with these challenges.^{72,74} In the era of increased interest in measuring functional health status, more exploration of the reasons for the divergence from a sequelae-based approach to measuring functional health status used in GBD versus an overall self-rating will be an important avenue of future research.

Crosswalks

Because of the diversity of data sources available, we estimated the statistical association between measurements taken using different case definitions, different assays, different survey items, or different ascertainment methods to a reference category. We used these statistical associations to crosswalk each type of data to the reference category. Crosswalks are a crucial dimension in the GBD analysis; in most cases we estimate these crosswalks from within DisMod-MR 2.1. If crosswalks were believed to vary substantially by age, sex, or location, these crosswalks were estimated outside of DisMod-MR 2.1 and adjusted data were used as inputs. Where data are sparse, the estimated crosswalks can shift substantially when new studies are identified that inform the crosswalk. With each iteration of the GBD, we have recognised the crucial nature of the crosswalk analysis for data processing. We believe that more research is needed in future iterations of GBD to standardise the approaches used for estimating crosswalks across disorders and risk factors, and for propagating uncertainty in crosswalks into the final results.

Data gaps

We have described the availability of data by cause and geography over different time periods, highlighting very large gaps in information availability. The data availability by country ranged from 6·1% in North Korea to 91·3% in the USA (methods appendix p 606). Among the top five causes of YLDs, data availability was rather poor. Lower back pain, iron-deficiency anaemia, major depressive disorder, other hearing loss, and neck pain all have data representativeness indices below 50%. Because of the broad overview provided of major data sources for disease

incidence and prevalence, GBD provides a framework to assist countries in prioritising the collection of new data to inform better monitoring of functional health status. By setting clear reference case definitions and data collection methods (methods appendix pp 4–7), GBD can also provide guidance on how to collect information most relevant to population health measurement.

Future directions for GBD

With each cycle of the GBD we expect, given the present interest in subnational assessments, to report increasingly granular results. These subnational findings will be of important national policy interest, but the discipline of examining the evidence for each community and modelling at the more granular level will, we believe, improve the quality of the national estimation as well. Additions to the cause list will continue to be driven by policy interest, such as the need to incorporate Zika virus, to split diabetes into separate estimates for type 1 and type 2, and by adding diseases to our cause list that are main contributors to large residual categories, such as other cardiovascular disease, other musculoskeletal disorders, and other neurological disorders. We plan to continue to expand our networks of topic-specific and country collaborators to enhance the quality of our estimates through their feedback and by enhancing the amount of data we can bring to bear on GBD estimation. Our country collaborations to generate subnational estimates have shown us that these efforts greatly enhance access to valuable data sources, particularly administrative datasets on inpatient and outpatient episodes and unpublished surveys.

Limitations

The GBD 2015 study has some key limitations. First, although we have sought to capture and include in our estimations many sources of uncertainty, we have not captured all sources. We have not been able to routinely capture uncertainty due to different model specifications in our results. We do not have the ability to reflect the uncertainty of data sources that exist but of which we are not aware. Subnational collaborations in China and Mexico revealed many data sources not captured in previous iterations of the GBD study. Inclusion of these data sources can lead to shifts in estimates that are outside the previously estimated 95% UIs.

Second, within DisMod-MR 2.1 we estimate the average association between different types of studies for reporting on a specific outcome such as the difference between 12-month recall and point prevalence in a survey. These estimated associations from within the DisMod-MR 2.1 likelihood estimation are used to adjust the data to a reference case definition or study design. These estimated adjustment factors are themselves uncertain, which increases the overall uncertainty in our estimation. More standardisation in data collection would be highly desirable and would reduce our dependence on the relatively challenging estimation process involved in crosswalks.

Third, for some disorders the data available to estimate excess mortality by age and sex and to capture how excess mortality changes with development status are very limited. There are probably many unpublished sources of information in some countries that could be usefully brought to bear on the challenge of estimating the age–sex–year–location levels of excess mortality.

Fourth, the microsimulation step of this study assumes that prevalence within an age–sex–location–year group of different sequelae is independent. Although this is clearly known not to be the case for some pairs of disorders, the independence assumption provides reasonable overall adjustments for comorbidity. Progress in incorporating dependent comorbidity has been difficult because information on the correlation structure of prevalence is extremely limited and only available for a minor fraction of all possible pairs of conditions in the GBD study.

Fifth, we estimated separate DisMod-MR 2.1 models for 1990, 1995, 2000, 2005, 2010, and 2015. Independent estimation of each time period implies that the uncertainty intervals for each period are also independent. Furthermore, compositional bias in the data available in different time periods might lead to spurious time trends. A more appealing strategy would be to simultaneously estimate the trends in incidence, excess mortality, and remission by age and sex consistent with all the available data for a geography on prevalence, incidence, remission, excess mortality, and cause-specific mortality by age and sex. DisMod-MR 2.1 does not allow for time-varying trends in incidence, excess mortality, and remission, but a prototype DisMod-MR AT (age and time) has been developed and is being tested. Allowing for all rates to change at different paces over age and time increases the number of parameters that need to be estimated by orders of magnitude.

Sixth, we have emphasised in the reporting of results the changes from calendar year 2005 to calendar year 2015, although we have estimated results for 1990, 1995, 2000, 2005, 2010, and 2015 and interpolated for the years in between. For some disorders, reporting the change from 2005 to 2015 (such as for Ebola virus disease) does not capture the major epidemic in west Africa in 2014. Likewise, disasters and wars are often concentrated in a single year; comparisons of any two years can provide misleading inferences about trends.

Seventh, for a few disorders (eg, tetanus, neonatal sepsis, rabies, and diphtheria) we estimate disease incidence from estimates of mortality and the inverse of the case-fatality rate estimated from available studies. When the case-fatality rate is very low, these estimates of incidence are highly sensitive to very small changes in the estimated case-fatality rate and have large uncertainty intervals.

Finally, although extraordinary efforts have gone into vetting the results by GBD researchers and the collaborator network, there are probably some findings that have not been scrutinised carefully enough. Making all results and underlying data available and having an increasing number of visualisation tools are effective

strategies that we will keep expanding to meet our goal of producing the highest quality global health data to our growing audience of policy makers, researchers, the media, and the general population.

Conclusion

The GBD studies seek to quantify the prevalence and incidence of the major sequelae for a comprehensive list of diseases and injuries; given the diversity of data sources, the range of biases in these sources, and the gaps in availability, this is a challenging task. Despite the limitations, the standardised and comprehensive approach of the GBD studies study provides useful insights. We believe users of the findings, including the wealth of country-specific and cause-specific detail available in the appendix, can usefully examine the broad trends for their country or subnational geography, benchmark against geographies at a similar level of development, and understand the strength or weakness of these estimates. Regular quantification of health is particularly important as the new health-related targets of the Sustainable Development Goals have broadened the health agenda throughout the world. At the same time, development and transformations in the risks to health experienced by different groups in the world are leading to some broad transformations in health. Everyone needs to have access to the timeliest, valid, reliable, and local information possible to enrich debates on how to accelerate health progress in all communities.

GBD 2015 Disease and Injury Incidence and Prevalence Collaborators

Theo Vos†, Christine Allen, Megha Arora, Ryan M Barber, Zulfiqar A Bhutta, Alexandria Brown, Austin Carter, Daniel C Casey, Fiona J Charlson, Alan Z Chen, Megan Coggeshall, Leslie Cornaby, Lalit Dandona, Daniel J Dicker, Tina Dilegge, Holly E Erskine, Alize J Ferrari, Christina Fitzmaurice, Tom Fleming, Mohammad H Forouzanfar, Nancy Fullman, Peter W Gething, Ellen M Goldberg, Nicholas Graetz, Juanita A Haagsma, Simon I Hay, Catherine O Johnson, Nicholas J Kasheba, Toana Kawashima, Laura Kemmer, Ibrahim A Khalil, Yohannes Kinfu, Hmwe H Kyu, Janni Leung, Xiaofeng Liang, Stephen S Lim, Alan D Lopez, Rafael Lozano, Laurie Marczak, George A Mensah, Ali H Mokdad, Mohsen Naghavi, Grant Nguyen, Elaine Nsoesie, Helen Olsen, David M Pigott, Christine Pinho, Zane Rankin, Nikolas Reinig, Joshua A Salomon, Logan Sandar, Alison Smith, Jeffrey Stanaway, Caitlyn Steiner, Stephanie Teeple, Bernadette A Thomas, Christopher Troeger, Joseph A Wagner, Haidong Wang, Valentine Wanga, Harvey A Whiteford, Leo Zoccker, Amanuel Alemu Abajobir*, Kalkidan Hassen Abate*, Cristiana Abbafati*, Kaja M Abbas*, Foad Abd-Allah*, Bijou Abraham*, Ibrahim Abubakar*, Laith J Abu-Raddad*, Niveen M E Abu-Rmeileh*, Ilana N Ackerman*, Akindele Olupelumi Adebisi*, Zafina Ademi*, Arsène Kouablan Adou*, Kossivi Agbenlenko Afanvi*, Emilie Elisabet Agardh*, Arnav Agarwal*, Aliasghar Ahmad Kiadaliri*, Hamid Ahmadi*, Oluremi N Ajala*, Rufus Olusola Akinyemi*, Nadia Akseer*, Ziyad Al-Aly*, Khurshid Alam*, Noore K M Alam*, Saleh Fahed Aldahri*, Miguel Angel Alegretti*, Zewdie Aderaw Alemu*, Lily T Alexander*, Samia Alhabib*, Raghib Ali*, Ala'a Alkerwi*, François Alla*, Peter Allebeck*, Rajaa Al-Raddadi*, Ubai Alsharif*, Khalid A Altirkawi*, Nelson Alvis-Guzman*, Azmeraw T Amare*, Alemayehu Amberbir*, Heresh Amini*, Walid Ammar*, Stephen Marc Amrock*, Hjalte H Andersen*, Gregory M Anderson*, Benjamin O Anderson*, Carl Abelardo T Antonio*, Atsed Fantahun Aregay*, Johan Årnlöv*, Al Artaman*, Hamid Asayesh*, Reza Assadi*, Suleman Atique*,

Euripide Frinel G Arthur Avokpaho*, Ashish Awasthi*, Beatriz Paulina Ayala Quintanilla*, Peter Azzopardi*, Umar Bacha*, Alaa Badawi*, Kalpana Balakrishnan*, Amitava Banerjee*, Aleksandra Barac*, Suzanne L Barker-Collo*, Till Bärnighausen*, Lars Barregard*, Lope H Barrero*, Arindam Basu*, Shahrzad Bazargan-Hejazi*, Ettore Beghi*, Brent Bell*, Michelle L Bell*, Derrick A Bennett*, Isabela M Bensenor*, Habib Benizian*, Adugnaw Berhane*, Eduardo Bernabé*, Balem Demtsu Betsu*, Addisu Shunu Beyene*, Neeraj Bhala*, Samir Bhatt*, Sibhatu Biadgilign*, Kelly Bienhoff*, Boris Bikbov*, Stan Biryukov*, Donal Bisanzio*, Espen Bjertness*, Jed Blome*, Rohan Borschmann*, Soufiane Boufous*, Michael Brainin*, Alexandra Brazinova*, Nicholas J K Breitborde*, Jonathan Brown*, Rachelle Buchbinder*, Geoffrey Colin Buckle*, Zahid A Butt*, Bianca Calabria*, Ismael Ricardo Campos-Nonato*, Julio Cesar Campuzano*, Hélène Carabin*, Rosario Cárdenas*, David O Carpenter*, Juan Jesus Carrero*, Carlos A Castañeda-Orjuela*, Jacqueline Castillo Rivas*, Ferrán Catalá-López*, Jung-Chen Chang*, Peggy Pei-Chia Chiang*, Chioma Ezinne Chibueze*, Vesper Hichilombwe Chisumpa*, Jee-Young Jasmine Choi*, Rajiv Chowdhury*, Hanne Christensen*, Devasahayam Jesudas Christopher*, Liliana G Ciobanu*, Massimo Cirillo*, Matthew M Coates*, Samantha M Colquhoun*, Cyrus Cooper*, Monica Cortinovis*, John A Crump*, Solomon Abrha Damtew*, Rakhi Dandona*, Farah Daoud*, Paul I Dargan*, José das Neves*, Gail Davey*, Adrian C Davis*, Diego De Leo*, Louisa Degenhardt*, Liana C Del Gobbo*, Robert P Dellavalle*, Kebede Deribe*, Amare Deribew*, Sarah Derrett*, Don C Des Jarlais*, Samath D Dharmaratne*, Preet K Dhillon*, Cesar Diaz-Torne*, Eric L Ding*, Tim R Driscoll*, Leilei Duan*, Manisha Dubey*, Bruce Bartholow Duncan*, Hedyeh Ebrahimi*, Richard G Ellenbogen*, Iqbal Elyazar*, Matthias Endres*, Aman Yesuf Endries*, Sergey Petrovich Ermakov*, Babak Eshtrati*, Kara Estep*, Talha A Farid*, Carla Sofia e Sa Farinha*, André Faro*, Maryam S Farvid*, Farshad Farzadfar*, Valery L Feigin*, David T Felson*, Seyed-Mohammad Fereshtehnejad*, Jefferson G Fernandes*, Joao C Fernandes*, Florian Fischer*, Joseph R A Fitchett*, Kyle Foreman*, F Gerry R Fowkes*, Jordan Fox*, Richard C Franklin*, Joseph Friedman*, Joseph Frostad*, Thomas Fürst*, Neal D Futran*, Belinda Gabbe*, Parthasarathi Ganguly*, Fortuné Gbètoho Gankpé*, Teshome Gebre*, Tsegaye Tewelde Gebrehiwot*, Amanuel Tesfay Gebremedhin*, Johanna M Geleijnse*, Bradford D Gessner*, Katherine B Gibney*, Ibrahim Abdelmageem Mohamed Ginawi*, Ababi Zergaw Giref*, Maurice Giroud*, Melkamu Dedefo Gishu*, Giorgia Giussani*, Elizabeth Glaser*, William W Godwin*, Hector Gomez-Dantes*, Philimon Gona*, Amador Goodridge*, Sameer Vali Gopalani*, Carolyn C Gotay*, Atsushi Goto*, Hebe N Gouda*, Rebecca Grainger*, Felix Greaves*, Francis Guillemin*, Yuming Guo*, Rahul Gupta*, Rajeev Gupta*, Vipin Gupta*, Reyna A Gutierrez*, Demewoz Haile*, Alemayehu Desalegne Hailu*, Gessesew Bugssa Hailu*, Yara A Halasa*, Randah Ribhi Hamadeh*, Samer Hamidi*, Mouhanad Hammami*, Jamie Hancock*, Alexis J Handal*, Graeme J Hankey*, Yuantao Hao*, Hilda L Harb*, Sivadasanpillai Harikrishnan*, Josep Maria Haro*, Rasmus Havmoeller*, Roderick J Hay*, Ileana Beatriz Heredia-Pi*, Pouria Heydarpour*, Hans W Hoek*, Masako Horino*, Nobuyuki Horita*, H Dean Hosgood*, Damian G Hoy*, Aung Soe Htet*, Hsiang Huang*, John J Huang*, Chantal Huynh*, Marissa Iannarone*, Kim Moesgaard Iburg*, Kaire Innos*, Manami Inoue*, Veena J Iyer*, Kathryn H Jacobsen*, Nader Jahanmehr*, Mihajlo B Jakovljevic*, Mehdi Javanbakht*, Sudha P Jayaraman*, Achala Upendra Jayatilake*, Sun Ha Jee*, Panniyammakal Jeemon*, Paul N Jensen*, Ying Jiang*, Tariku Jibat*, Aida Jimenez-Corona*, Ye Jin*, Jost B Jonas*, Zubair Kabir*, Yogeshwar Kalkonde*, Ritul Kamal*, Haidong Kan*, André Karch*, Corine Kakizi Karema*, Chante Karimkhani*, Amir Kasaieian*, Anil Kaul*, Norito Kawakami*, Peter Njenga Keiyoro*, Andrew Haddon Kemp*, Andre Keren*, Chandrasekharan Nair Kesavachandran*, Yousef Saleh Khader*, Abdur Rahman Khan*, Ejaz Ahmad Khan*, Young-Ho Khang*,

- Sahil Khera*, Tawfik Ahmed Muthafer Khoja*, Jagdish Khubchandani*, Christian Kielsing*, Pauline Kim*, Cho-il Kim*, Daniel Kim*, Yun Jin Kim*, Niranjan Kissoon*, Luke D Knibbs*, Ann Kristin Knudsen*, Yoshihiro Kokubo*, Dhaval Kolte*, Jacek A Kopec*, Soewarta Kosen*, Georgios A Kotsakis*, Parvaiz A Koul*, Ai Koyanagi*, Michael Kravchenko*, Barthelemy Kuate Defo*, Burcu Kucuk Bicer*, Andreas A Kudom*, Ernst J Kuipers*, G Anil Kumar*, Michael Kutz*, Gene F Kwan*, Aparna Lal*, Ratilal Laloo*, Tea Lallukka*, Hilton Lam*, Jennifer O Lam*, Sinead M Langan*, Anders Larsson*, Pablo M Lavados*, Janet L Leasher*, James Leigh*, Ricky Leung*, Miriam Levi*, Yichong Li*, Yongmei Li*, Juan Liang*, Shiwei Liu*, Yang Liu*, Belinda K Lloyd*, Warren D Lo*, Giancarlo Logroscino*, Katharine J Looker*, Paulo A Lotufo*, Raimundas Lunevicius*, Ronan A Lyons*, Mark T Mackay*, Mohammed Magdy Abd El Razek*, Mahdi Mahdavi*, Marek Majdan*, Azeem Majeed*, Reza Malekzadeh*, Wagner Marcenes*, David Joel Margolis*, Jose Martinez-Raga*, Felix Masiye*, João Massano*, Stephen Theodore McGarvey*, John J McGrath*, Martin McKee*, Brian J McMahon*, Peter A Meaney*, Alem Mehari*, Fabiola Mejia-Rodriguez*, Alemayehu B Mekonnen*, Yohannes Adama Melaku*, Peter Memiah*, Ziad A Memish*, Walter Mendoza*, Atte Meretoja*, Tuomo J Meretoja*, Francis Apolinary Mhimbira*, Anoushka Millar*, Ted R Miller*, Edward J Mills*, Mojde Mirarefin*, Philip B Mitchell*, Charles N Mock*, Alireza Mohammadi*, Shafiu Mohammed*, Lorenzo Monasta*, Julio Cesar Montañez Hernandez*, Marcella Montico*, Meghan D Mooney*, Maziar Moradi-Lakeh*, Lidia Morawska*, Ulrich O Mueller*, Erin Mullany*, John Everett Mumford*, Michele E Murdoch*, Jean B Nachega*, Gabriele Nagel*, Aliya Naheed*, Luigi Naldi*, Vinay Nangia*, John N Newton*, Marie Ng*, Frida Namnyak Ngalesoni*, Quyen Le Nguyen*, Muhammad Imran Nisar*, Patrick Martial Nkamedjie Pete*, Joan M Nolla*, Ole F Norheim*, Rosana E Norman*, Bo Norrving*, Bruno P Nunes*, Felix Akpojene Ogbo*, In-Hwan Oh*, Takayoshi Ohkubo*, Pedro R Olivares*, Bolajoko Olubukunola Olusanya*, Jacob Olusegun Olusanya*, Alberto Ortiz*, Majdi Osman*, Erika Ota*, Mahesh PA*, Eun-Kee Park*, Mahboubeh Parsaiean*, Valéria Maria de Azeredo Passos*, Angel J Paternina Caicedo*, Scott B Patten*, George C Patton*, David M Pereira*, Rogelio Perez-Padilla*, Norberto Perico*, Konrad Pesudovs*, Max Petzold*, Michael Robert Phillips*, Frédéric B Piel*, Julian David Pillay*, Farhad Pishgar*, Dietrich Plass*, James A Platts-Mills*, Suzanne Polinder*, Constance D Pond*, Svetlana Popova*, Richie G Poulton*, Farshad Pourmalek*, Dorairaj Prabhakaran*, Noela M Prasad*, Mostafa Qorbani*, Rynaz H S Rabiee*, Amir Radfar*, Anwar Rafay*, Kazem Rahimi*, Vafa Rahimi-Movaghar*, Mahfuzar Rahman*, Mohammad Hifz Ur Rahman*, Sajjad Ur Rahman*, Rajesh Kumar Rai*, Sasa Rajsic*, Usha Ram*, Puja Rao*, Amany H Refaat*, Marissa B Reitsma*, Giuseppe Remuzzi*, Serge Resnikoff*, Alex Reynolds*, Antonio L Ribeiro*, Maria Jesus Rios Blancas*, Hirbo Shore Roba*, David Rojas-Rueda*, Luca Ronfani*, Gholamreza Roshandel*, Gregory A Roth*, Dietrich Rothenbacher*, Ambuj Roy*, Rajesh Sagar*, Ramesh Sahathevan*, Juan R Sanabria*, Maria Dolores Sanchez-Niño*, Itamar S Santos*, João Vasco Santos*, Rodrigo Sarmiento-Suarez*, Benn Sartorius*, Maheswar Satpathy*, Miloje Savic*, Monika Sawhney*, Michael P Schaub*, Maria Inês Schmidt*, Ione J C Schneider*, Ben Schöttker*, David C Schwebel*, James G Scott*, Soraya Seedat*, Sadaf G Sepanlou*, Edson E Servan-Mori*, Katya A Shackelford*, Amira Shaheen*, Masood Ali Shaikh*, Rajesh Sharma*, Upasana Sharma*, Jiabin Shen*, Donald S Shepard*, Kevin N Sheth*, Kenji Shibuya*, Min-Jeong Shin*, Rahman Shiri*, Ivy Shiue*, Mark G Shrim*, Inga Dora Sigfusdottir*, Diego Augusto Santos Silva*, Dayane Gabriele Alves Silveira*, Abhishek Singh*, Jasvinder A Singh*, Om Prakash Singh*, Prashant Kumar Singh*, Anna Sivonda*, Vegard Skirbekk*, Jens Christoffer Skogen*, Amber Sligar*, Karen Sliwa*, Michael Soljak*, Kjetil Søreide*, Reed J D Sorensen*, Joan B Soriano*, Luciano A Sposato*, Chandrashekar T Sreeramareddy*, Vasiliki Stathopoulou*, Nicholas Steel*, Dan J Stein*, Timothy J Steiner*, Sabine Steinke*, Lars Stovner*, Konstantinos Stroumpoulis*, Bruno F Sunguya*, Patrick Sur*, Soumya Swaminathan*, Bryan L Sykes*, Cassandra E I Szoek*, Rafael Tabarés-Seisdedos*, Jukka S Takala*, Nikhil Tandon*, David Tanne*, Mohammad Tavakkoli*, Bineyam Teye*, Hugh R Taylor*, Braden J Te Ao*, Bemnet Amare Tedla*, Abdullah Sulieman Terkawi*, Alan J Thomson*, Andrew L Thorne-Lyman*, Amanda G Thrift*, George D Thurston*, Ruoyan Tobe-Gai*, Marcello Tonelli*, Roman Topor-Madry*, Fotis Topouzis*, Bach Xuan Tran*, Thomas Truelsen*, Zacharie Tsala Dimbuene*, Miltiadis Tsimbaris*, Abera Kenay Tura*, Emin Murat Tuzcu*, Stefanos Tyrovolas*, Kingsley N Ukwaja*, Eduardo A Undurraga*, Chigozie Jesse Uneke*, Olalekan A Uthman*, Coen H van Gool*, Yuri Y Varakin*, Tommi Vasankari*, Narayanaswamy Venketasubramanian*, Raj Kumar Verma*, Francesco S Violante*, Sergey K Vladimirov*, Vasilii Victorovich Vlassov*, Stein Emil Vollset*, Gregory R Wagner*, Stephen G Waller*, Linhong Wang*, David A Watkins*, Scott Weichenath*, Elisabete Weiderpass*, Robert G Weintraub*, Andrea Werdecker*, Ronny Westerman*, Richard A White*, Hywel C Williams*, Charles Shey Wiyong*, Charles D A Wolfe*, Sungho Won*, Rachel Woodbrook*, Mamo Wubshet*, Denis Xavier*, Gelin Xu*, Ajit Kumar Yadav*, Lijing L Yan*, Yuichiro Yano*, Mehdi Yaseri*, Pengpeng Ye*, Henock Gebremedhin Yebo*, Paul Yip*, Naohiro Yonemoto*, Seok-Jun Yoon*, Mustafa Z Younis*, Chuanhua Yu*, Zoubida Zaidi*, Maysaa El Sayed Zaki*, Hajo Zeeb*, Maigeng Zhou*, Sanjay Zodpey*, Liesl Joanna Zuhlke*, Christopher J L Murray.
- *Authors listed alphabetically. †Corresponding author.
- Affiliations**
 Institute for Health Metrics and Evaluation (Prof T Vos PhD, C Allen BA, M Arora BSA, R M Barber BS, A Brown MA, A Carter BS, D C Casey BA, F J Charlson PhD, A Z Chen BS, M Coggeshall BA, L Cornaby BS, Prof L Dandona MD, D J Dicker BS, T Dilegge BS, H E Erskine PhD, A J Ferrari PhD, C Fitzmaurice MD, T Fleming BS, M H Forouzanfar MD, N Fullman MPH, E M Goldberg BS, N Graetz MPH, J A Haagsma PhD, Prof S I Hay DSc, C O Johnson PhD, N J Kashebaum MD, T Kawashima MS, L Kemmer PhD, I A Khalil MD, H H Kyu PhD, J Leung PhD, Prof S S Lim PhD, Prof A D Lopez PhD, L Marczak PhD, Prof A H Mokdad PhD, Prof M Naghavi PhD, G Nguyen MPH, E Nsoesie PhD, H Olsen MAIS, D M Pigott DPhil, C Pinho BA, Z Rankin BS, N Reinig BS, L Sandar BS, A Smith BA, J Stanaway PhD, C Steiner MPH, S Teplee BA, B A Thomas MD, C Troeger MPH, J A Wagner BS, H Wang PhD, V Wang MS, Prof H A Whiteford PhD, L Zoeckler BA, L T Alexander BA, G M Anderson MSEE, B Bell MLIS, K Bienhoff MA, S Biryukov BS, J Blore PhD, J Brown MAIS, M M Coates MPH, F Daoud BS, K Estep MPA, K Foreman PhD, J Fox BS, J Friedman BA, J Frostad MPH, W W Godwin BS, J Hancock MLS, C Huynh BA, M Iannarone MSc, P Kim BA, M Kutz BS, F Masiye PhD, A Millar BA, M Mirarefin MPH, M D Mooney BS, M Moradi-Lakeh MD, E Mullany BA, J E Mumford BA, M Ng PhD, P Rao MPH, M B Reitsma BS, A Reynolds BA, G A Roth MD, K A Shackelford BA, A Sivonda MHA, A Sligar MPH, R J D Sorensen MPH, P Sur BA, Prof S E Vollset DrPH, R Woodbrook MLIS/MA, Prof M Zhou PhD, Prof C J L Murray DPhil, Harborview/UW Medicine (R G Ellenbogen MD), School of Dentistry (G A Kotsakis DDS), Harborview Injury Prevention and Research Center (C N Mock PhD), University of Washington, Seattle, WA, USA (Prof B O Anderson MD, N D Futran MD, P N Jensen PhD, D A Watkins MD); Centre of Excellence in Women and Child Health (Prof Z A Bhutta PhD), Aga Khan University, Karachi, Pakistan (M I Nisar MSc); Centre for Global Child Health, The Hospital for Sick Children, Toronto, ON, Canada (Prof Z A Bhutta PhD, N Akseer MSc); School of Public Health (F J Charlson PhD, H E Erskine PhD, A J Ferrari PhD, J Leung PhD, Prof H A Whiteford PhD, A A Abajobir MPH, L D Knibbs PhD), School of Dentistry (Prof R Laloo PhD), Centre for Clinical Research (J G Scott PhD), The University of Queensland, Brisbane, QLD, Australia (N K M Alam MPH, H N Gouda PhD, Y Guo PhD, Prof J J McGrath MD); Queensland Centre for Mental Health Research, Brisbane, QLD, Australia (F J Charlson PhD, H E Erskine PhD, A J Ferrari PhD, J Leung PhD, Prof H A Whiteford PhD); Centre for Control of Chronic

Conditions (P Jeemon PhD), Public Health Foundation of India, New Delhi, India (Prof L Dandona MD, R Dandona PhD, G A Kumar PhD); Department of Zoology (P W Gething PhD), Nuffield Department of Medicine (D Bisanzio PhD, A Deribew PhD), NIHR Musculoskeletal Biomedical Research Centre (Prof C Cooper FMedSci), Oxford Big Data Institute, Li Ka Shing Centre for Health Information and Discovery (Prof S I Hay DSc), University of Oxford, Oxford, UK (R Ali FRCP, D A Bennett PhD, K Rahimi DM); Centre for Research & Action in Public Health, Faculty of Health, University of Canberra, Canberra, ACT, Australia (Y Kinfu PhD); National Center for Chronic and Noncommunicable Disease Control and Prevention (L Duan MD, Y Li MPH, S Liu PhD, Y Jin MS, Prof L Wang MD, P Ye MPH, Prof M Zhou PhD), Chinese Center for Disease Control and Prevention (Prof X Liang MD), Beijing, China; Melbourne School of Population and Global Health (Prof A D Lopez PhD), Department of Paediatrics (P Azzopardi MEpi), The Peter Doherty Institute for Infection and Immunity (K B Gibney FRACP), Department of Medicine (A Meretoja PhD), Murdoch Childrens Research Institute (K Alam PhD, P Azzopardi MEpi, R Borschmann PhD, S M Colquhoun PhD, Prof G C Patton MD, R G Weintraub MBBS), Institute of Health and Ageing (Prof C E I Szoek PhD), The University of Melbourne, Melbourne, VIC, Australia (Z Ademi PhD, K Alam PhD, R Borschmann PhD, S M Colquhoun PhD, Prof H R Taylor AC, R G Weintraub MBBS); National Institute of Public Health, Cuernavaca, Mexico (R Lozano MD, I R Campos-Nonato PhD, J C Campuzano PhD, H Gomez-Dantes MSc, I B Heredia-Pi PhD, F Mejia-Rodriguez MD, J C Montañez Hernandez MSc, M J Rios Blancas MPH, Prof E E Servan-Mori MSc); Center for Translation Research and Implementation Science, National Heart, Lung, and Blood Institute, National Institutes of Health, Bethesda, MD, USA (G A Mensah MD); Department of Global Health and Population (Prof J A Salomon PhD), Department of Nutrition (A L Thorne-Lyman ScD), Harvard T H Chan School of Public Health (O N Ajala MD, Prof T Bärnighausen MD, I R Campos-Nonato PhD, E L Ding ScD, M S Farvid PhD, G R Wagner MD), Harvard Medical School (M Osman MD, M G Shrimme MD), Harvard University, Boston, MA, USA (J R A Fitchett MD); Jimma University, Jimma, Ethiopia (K H Abate MS, T T Gebrehiwot MPH, A T Gebremedhin MPH); La Sapienza, University of Rome, Rome, Italy (C Abbafati PhD); Virginia Tech, Blacksburg, VA, USA (Prof K M Abbas PhD); Department of Neurology, Cairo University, Cairo, Egypt (Prof F Abd-Allah MD); NMSM Government College Kalpetta, Kerala, India (Prof B Abraham MPhil); Institute for Global Health (Prof I Abubakar PhD), Farr Institute of Health Informatics Research (A Banerjee DPhil), Department of Epidemiology and Public Health (H Benzian PhD), University College London, London, UK; Infectious Disease Epidemiology Group, Weill Cornell Medical College in Qatar, Doha, Qatar (L J Abu-Raddad PhD); Institute of Community and Public Health, Birzeit University, Ramallah, Palestine (N M Abu-Rmeileh PhD); Department of Epidemiology and Preventive Medicine, School of Public Health and Preventive Medicine (I N Ackerman PhD, Prof R Buchbinder PhD), School of Public Health and Preventive Medicine (Prof B Gabbe PhD), Department of Medicine, School of Clinical Sciences at Monash Health (Prof A G Thrift PhD), Monash University, Melbourne, VIC, Australia; College of Medicine (A O Adebiyi MD), University of Ibadan, Ibadan, Nigeria (R O Akinyemi PhD); University College Hospital, Ibadan, Nigeria (A O Adebiyi MD); University of Basel, Basel, Switzerland (Z Ademi PhD, T Fürst PhD); Association Ivoirienne pour le Bien-Être Familial, Abidjan, Côte d'Ivoire (A K Adou MD); Direction du District Sanitaire de Haho, Notse, Togo (K A Afanvi MD); Faculté des Sciences de Santé, Université de Lomé, Lomé, Togo (K A Afanvi MD); Institution of Public Health Sciences, Stockholm, Sweden (E E Agardh PhD); Dalla Lana School of Public Health (N Akseer MSc), Department of Nutritional Sciences, Faculty of Medicine (A Badawi PhD), Centre for Addiction and Mental Health (S Popova PhD), University of Toronto, Toronto, ON, Canada (A Agarwal BHSc); McMaster University, Hamilton, ON, Canada (A Agarwal BHSc); Department of Clinical Sciences Lund, Orthopedics, Clinical Epidemiology Unit (A Ahmad Kiadaliri PhD), Skane University Hospital, Department of Clinical Sciences Lund (Prof B Norrving PhD), Lund University, Lund,

Sweden; Health Services Management Research Center, Institute for Futures Studies in Health, Kerman University of Medical Sciences, Kerman, Iran (A Ahmad Kiadaliri PhD); Ophthalmic Research Center (H Ahmadi MD, M Yaseri PhD), School of Public Health (N Jahanmehr PhD), Shahid Beheshti University of Medical Sciences, Tehran, Iran; Department of Ophthalmology, Labbafinejad Medical Center, Tehran, Iran (H Ahmadi MD); University of Pittsburgh Medical Center, McKeesport, PA, USA (O N Ajala MD); Newcastle University, Newcastle upon Tyne, UK (R O Akinyemi PhD); Washington University in Saint Louis, St Louis, MO, USA (Z Al-Aly MD); Sydney School of Public Health (Prof T R Driscoll PhD), The University of Sydney, Sydney, NSW, Australia (K Alam PhD, Prof A H Kemp PhD, J Leigh PhD, A B Mekonnen MS); Queensland Health, Brisbane, QLD, Australia (N K M Alam MPH); King Saud University, Riyadh, Saudi Arabia (S F Aldahri MD, K A Altirkawi MD); King Fahad Medical City, Riyadh, Saudi Arabia (S F Aldahri MD); Department of Preventive and Social Medicine, Faculty of Medicine, University of the Republic, Montevideo, Uruguay (M A Alegratti MD); Debre Markos University, Debre Markos, Ethiopia (Z A Alemu MPH); King Abdullah Bin Abdulaziz University Hospital, Riyadh, Saudi Arabia (S Alhabib PhD); Luxembourg Institute of Health (LIH), Strassen, Luxembourg (A Alkerwi PhD); School of Public Health, University of Lorraine, Nancy, France (Prof F Alla PhD, F Guillemin PhD); Department of Public Health Sciences (P Allebeck PhD, R H S Rabiee MPH), Department of Clinical Science, Intervention and Technology (Prof J J Carrero PhD), Department of Neurobiology, Care Sciences and Society (NVS) (S M Ferestehnejad PhD), Department of Medical Epidemiology and Biostatistics (E Weiderpass PhD), Karolinska Institutet, Stockholm, Sweden (R Havmoeller PhD); Ministry of Health, Jeddah, Saudi Arabia (R Al-Raddadi PhD); Charité Universitätsmedizin, Berlin, Germany (U Alsharif MPH); Universidad de Cartagena, Cartagena de Indias, Colombia (Prof N Alvis-Guzman PhD); School of Medicine (A T Amare MPH, Y A Melaku MPH), University of Adelaide, Adelaide, SA, Australia (L G Ciobanu MS); College of Medicine and Health Sciences, Bahir Dar University, Bahir Dar, Ethiopia (A T Amare MPH); Dignitas International, Zomba, Malawi (A Amberbir PhD); Environmental Health Research Center, Kurdistan University of Medical Sciences, Sanandaj, Iran (H Amini MSPH); Department of Epidemiology and Public Health (H Amini MSPH, T Fürst PhD), Swiss Tropical and Public Health Institute, Basel, Switzerland (C K Karema MSc); Ministry of Public Health, Beirut, Lebanon (W Ammar PhD, H L Harb MPH); Oregon Health & Science University, Portland, OR, USA (S M Amrock MD); Center for Sensory-Motor Interaction, Department of Health Science and Technology, Faculty of Medicine, Aalborg University, Aalborg, Denmark (H H Andersen MSc); Department of Health Policy and Administration, College of Public Health, University of the Philippines Manila, Manila, Philippines (C A T Antonio MD); School of Public Health (Y A Melaku MPH), Mekelle University, Mekelle, Ethiopia (A F Aregay MS, B D Betsu MS, G B Hailu MSc, H G Yebo MS); Department of Medical Sciences, Uppsala University, Uppsala, Sweden (Prof J Ärnlov PhD, Prof A Larsson PhD); Dalarna University, Falun, Sweden (Prof J Ärnlov PhD); Consultant, Windsor, ON, Canada (A Artaman PhD); Department of Medical Emergency, School of Paramedic, Qom University of Medical Sciences, Qom, Iran (H Asayesh PhD); Mashhad University of Medical Sciences, Mashhad, Iran (R Assadi PhD); Graduate Institute of Biomedical Informatics, Taipei Medical University, Taipei, Taiwan (S Atique MS); Institut de Recherche Clinique du Bénin, Cotonou, Benin Republic (E F G A Avokpaho MPH); Laboratoire d'Etudes et de Recherche-Action en Santé (LERAS Afrique), Parakou, Benin Republic (E F G A Avokpaho MPH); Sanjay Gandhi Postgraduate Institute of Medical Sciences, Lucknow, India (A Awasthi MSc); The Judith Lumley Centre for Mother, Infant and Family Health Research, La Trobe University, Melbourne, VIC, Australia (B P Ayala Quintanilla PhD); Peruvian National Institute of Health, Lima, Peru (B P Ayala Quintanilla PhD); Wardliparingga Aboriginal Research Unit, South Australian Health and Medical Research Institute, Adelaide, SA, Australia (P Azzopardi MEpi); School of Health Sciences, University of Management and Technology, Lahore, Pakistan (U Bacha PhD); Public Health Agency of Canada, Toronto, ON, Canada (A Badawi PhD);

- Department of Environmental Health Engineering, Sri Ramachandra University, Chennai, India (K Balakrishnan PhD); Faculty of Medicine, University of Belgrade, Belgrade, Serbia (A Barac PhD); School of Psychology, University of Auckland, Auckland, New Zealand (S L Barker-Collo PhD); Africa Health Research Institute, Mtubatuba, South Africa (Prof T Bärnighausen MD); Institute of Public Health, Heidelberg University, Heidelberg, Germany (Prof T Bärnighausen MD, S Mohammed PhD); Department of Occupational and Environmental Health (Prof L Barregard MD), Health Metrics Unit (Prof M Petzold PhD), University of Gothenburg, Gothenburg, Sweden; Department of Industrial Engineering, School of Engineering, Pontificia Universidad Javeriana, Bogotá, Colombia (L H Barrero ScD); School of Health Sciences, University of Canterbury, Christchurch, New Zealand (A Basu PhD); College of Medicine, Charles R Drew University of Medicine and Science, Los Angeles, CA, USA (Prof S Bazargan-Hejazi PhD); David Geffen School of Medicine, University of California, Los Angeles, Los Angeles, CA, USA (Prof S Bazargan-Hejazi PhD); Kermanshah University of Medical Science, Kermanshah, Iran (Prof S Bazargan-Hejazi PhD); IRCCS-Istituto di Ricerche Farmacologiche Mario Negri, Milan, Italy (E Beghi MD); School of Medicine (K N Sheth MD), Yale University, New Haven, CT, USA (Prof M L Bell PhD, J J Huang MD); Internal Medicine Department (Prof I S Santos PhD), University of São Paulo, São Paulo, Brazil (I M Bensenor PhD, Prof P A Lotufo DrPH); Department of Epidemiology and Health Promotion, College of Dentistry (H Benzan PhD), New York University, New York, NY, USA; Debre Berhane University, Debre Berhan, Ethiopia (A Berhane PhD); Division of Health and Social Care Research (Prof C D Wolfe MD), King's College London, London, UK (E Bernabé PhD, Prof R J Hay DM); College of Health and Medical Sciences (H S Roba MPH), Haramaya University, Harar, Ethiopia (A S Beyene MPH); Queen Elizabeth Hospital Birmingham, Birmingham, UK (N Bhala DPhil); University of Otago Medical School, Wellington, New Zealand (N Bhala DPhil); Department of Infectious Disease Epidemiology (T Fürst PhD), Department of Epidemiology and Biostatistics (F B Piel PhD), Division of Brain Sciences (Prof T J Steiner PhD), Imperial College London, London, UK (S Bhatt DPhil, F Greaves PhD, Prof A Majeed MD, M Soljak PhD); Independent Public Health Consultants, Addis Ababa, Ethiopia (S Biadgilign MPH); Department of Nephrology Issues of Transplanted Kidney, Academician V I Shumakov Federal Research Center of Transplantation and Artificial Organs, Moscow, Russia (B Bikbov MD); Department of Community Medicine (Prof E Bjertness PhD), University of Oslo, Oslo, Norway (A S Htet MPH); Transport and Road Safety (TARS) Research (S Boufous PhD), National Drug and Alcohol Research Centre (Prof L Degenhardt PhD), Brien Holden Vision Institute (Prof S Resnikoff MD), University of New South Wales, Kensington, NSW, Australia (B Calabria PhD, Prof P B Mitchell MD); Danube-University Krems, Krems, Austria (Prof M Brainin PhD); Faculty of Health Sciences and Social Work, Department of Public Health, Trnava University, Trnava, Slovakia (A Brazinova PhD, M Majdan PhD); International Neurotrauma Research Organization, Vienna, Austria (A Brazinova PhD); Departments of Pediatrics and Neurology (W D Lo MD), College of Medicine (J Shen PhD), The Ohio State University, Columbus, OH, USA (Prof N J K Breitborde PhD); Monash Department of Clinical Epidemiology, Cabrini Institute, Melbourne, VIC, Australia (Prof R Buchbinder PhD); University of California, San Francisco, San Francisco, CA, USA (G C Buckle MD); Al Shifa Trust Eye Hospital, Rawalpindi, Pakistan (Z A Butt PhD); National Centre for Epidemiology and Population Health, Australian National University, Canberra, ACT, Australia (B Calabria PhD, A Lal PhD); Department of Biostatistics and Epidemiology, University of Oklahoma Health Sciences Center, Oklahoma City, OK, USA (H Carabin PhD); Metropolitan Autonomous University, Mexico City, Mexico (R Cárdenas ScD); University at Albany, Rensselaer, NY, USA (Prof D O Carpenter MD); Colombian National Health Observatory, Instituto Nacional de Salud, Bogotá, Colombia (C A Castañeda-Orjuela MSc); Epidemiology and Public Health Evaluation Group, Public Health Department, Universidad Nacional de Colombia, Bogotá, Colombia (C A Castañeda-Orjuela MSc); Caja Costarricense de Seguro Social, San Jose, Costa Rica (Prof J Castillo Rivas MPH); Universidad de Costa Rica, San Pedro, Montes de Oca, Costa Rica (Prof J Castillo Rivas MPH); Department of Medicine, University of Valencia/INCLIVA Health Research Institute and CIBERSAM, Valencia, Spain (F Catalá-López PhD); Clinical Epidemiology Program, Ottawa Hospital Research Institute, Ottawa, ON, Canada (F Catalá-López PhD); School of Nursing, College of Medicine, National Taiwan University, Taipei, Taiwan (Prof J Chang PhD); Clinical Governance Unit, Gold Coast Health, Southport, QLD, Australia (P P Chiang PhD); National Center for Child Health and Development, Setagaya, Japan (C E Chibueze PhD); University of Zambia, Lusaka, Zambia (V H Chisumpa MPhil, F Masiye PhD); University of Witwatersrand, Johannesburg, South Africa (V H Chisumpa MPhil); Seoul National University Medical Library, Seoul, South Korea (J J Choi PhD); Department of Public Health and Primary Care, University of Cambridge, Cambridge, UK (R Chowdhury PhD); Bispebjerg University Hospital, Copenhagen, Denmark (Prof H Christensen DMSc); Christian Medical College, Vellore, India (Prof D J Christopher MD); University of Salerno, Baronissi, Italy (Prof M Cirillo MD); MRC Lifecourse Epidemiology Unit, University of Southampton, Southampton, UK (Prof C Cooper FMedSci); NIHR Biomedical Research Centre, University of Southampton and University Hospital Southampton NHS Foundation Trust, Southampton, UK (Prof C Cooper FMedSci); IRCCS-Istituto di Ricerche Farmacologiche Mario Negri, Bergamo, Italy (M Cortinovis Biotech D, G Giussani Biol, D N Perico MD, Prof G Remuzzi MD,); Centre for International Health, Dunedin School of Medicine (Prof J A Crump MD), Injury Prevention Research Unit, Department of Preventive and Social Medicine, Dunedin School of Medicine (Prof S Derrett PhD), University of Otago, Dunedin, New Zealand (Prof R G Poulton PhD); Wolaita Sodo University, Wolaita Sodo, Ethiopia (S A Damtew MPH); School of Public Health (K Deribe MPH, A D Hailu MPH), Addis Ababa University, Addis Ababa, Ethiopia (S A Damtew MPH, A Z Giref PhD, D Haile MPH, T Jibat MS, B Taye PhD); Guy's and St Thomas' NHS Foundation Trust, London, UK (Prof P I Dargan FRCP); i3S - Instituto de Investigação e Inovação em Saúde and INEB - Instituto de Engenharia Biomédica (J das Neves PhD), Faculty of Medicine (J Massano MD, J V Santos BHLthSc); University of Porto, Porto, Portugal; Wellcome Trust Brighton & Sussex Centre for Global Health Research, Brighton, UK (Prof G Davey MD); Public Health England, London, UK (Prof A C Davis PhD, F Greaves PhD, Prof J N Newton FRCP, Prof N Steel PhD); Griffith University, Brisbane, QLD, Australia (Prof D De Leo DSc); Stanford University, Stanford, CA, USA (L C Del Gobbo PhD); University of Colorado School of Medicine and the Colorado School of Public Health, Aurora, CO, USA (R P Dellavalle MD); Brighton and Sussex Medical School, Brighton, UK (K Deribe MPH); KEMRI-Wellcome Trust Research Programme, Kilifi, Kenya (A Deribew PhD); Mount Sinai Beth Israel, New York, NY, USA (Prof D C Des Jarlais PhD); Icahn School of Medicine at Mount Sinai, New York, NY, USA (Prof D C Des Jarlais PhD); Department of Community Medicine, Faculty of Medicine, University of Peradeniya, Peradeniya, Sri Lanka (S D Dharmaratne MD); Centre for Control of Chronic Conditions (P Jeemon PhD), Public Health Foundation of India, Gurgaon, India (P K Dhillon PhD, P Ganguly MD, Prof S Zodpey PhD); Hospital de la Santa Creu i Sant Pau, Barcelona, Spain (C Diaz-Torné MD); International Institute for Population Sciences, Mumbai, India (M Dubey MPhil, M H U Rahman MPhil, Prof U Ram PhD, A Singh PhD, R K Verma MPhil, A K Yadav MPhil); Federal University of Rio Grande do Sul, Porto Alegre, Brazil (B B Duncan PhD, C Kielsing MD, Prof M I Schmidt MD); University of North Carolina, Chapel Hill, NC, USA (B B Duncan PhD); Non-communicable Diseases Research Center, Endocrinology and Metabolism Population Sciences Institute (H Ebrahimi MD, F Pishgar MD, F Farzadfar MD, A Kasaieian PhD, M Parsaeian PhD), Liver and Pancreaticobiliary Diseases Research Center, Digestive Disease Research Institute, Shariati Hospital (H Ebrahimi MD), Multiple Sclerosis Research Center, Neuroscience Institute (P Heydarpour MD), Hematology-Oncology and Stem Cell Transplantation Research Center (A Kasaieian PhD), Digestive Diseases Research Institute (Prof R Malekzadeh MD, G Roshandel PhD, S G Sepanlou PhD), Department of Epidemiology and Biostatistics, School of Public Health (M Parsaeian PhD), Uro-Oncology Research Center (F Pishgar MD), Sina Trauma and Surgery Research Center (Prof V Rahimi-Movaghar MD), Tehran University of Medical Sciences, Tehran, Iran (M Yaseri PhD);

Eijkman-Oxford Clinical Research Unit, Jakarta, Indonesia (I Elyazar PhD); Charité University Medicine Berlin, Berlin, Germany (Prof M Endres MD); Arba Minch University, Arba Minch, Ethiopia (A Y Endries MPH); The Institute of Social and Economic Studies of Population, Russian Academy of Sciences, Moscow, Russia (Prof S P Ermakov DSc); Federal Research Institute for Health Organization and Informatics, Ministry of Health of the Russian Federation, Moscow, Russia (Prof S P Ermakov DSc); Ministry of Health and Medical Education, Tehran, Iran (B Eshtrati PhD); Arak University of Medical Sciences, Arak, Iran (B Eshtrati PhD); University of Louisville, Louisville, KY, USA (T A Farid MD, A R Khan MD); DGS Directorate General of Health, Lisboa, Portugal (C S E S Farinha MSc); Universidade Aberta, Lisboa, Portugal (C S E S Farinha MSc); Federal University of Sergipe, Aracaju, Brazil (Prof A Faro PhD); Harvard/MGH Center on Genomics, Vulnerable Populations, and Health Disparities, Mongan Institute for Health Policy, Massachusetts General Hospital, Boston, MA, USA (M S Farvid PhD); National Institute for Stroke and Applied Neurosciences (V L Feigin PhD), Auckland University of Technology, Auckland, New Zealand (B J Te Ao MPH); School of Medicine (G F Kwan MD), Boston University, Boston, MA, USA (Prof D T Felson MD); Institute of Education and Sciences, German Hospital Oswaldo Cruz, São Paulo, Brazil (Prof J G Fernandes PhD); Centre for Experimental Medicine & Rheumatology, William Harvey Research Institute, Barts and The London School of Medicine & Dentistry, Queen Mary University of London, London, UK (J C Fernandes PhD); Bielefeld University, Bielefeld, Germany (F Fischer MPH); Alzheimer Scotland Dementia Research Centre (I Shiue PhD), University of Edinburgh, Edinburgh, UK (Prof F G R Fowkes PhD); James Cook University, Townsville, QLD, Australia (R C Franklin PhD); Department of Infectious Disease Epidemiology (T Fürst PhD), Department of Epidemiology and Biostatistics (F B Piel PhD), Division of Brain Sciences (Prof T J Steiner PhD), Imperial College London, London, UK (F Greaves PhD, Prof A Majeed MD, M Soljak PhD); Indian Institute of Public Health Gandhinagar, Ahmedabad, India (P Ganguly MD, V J Iyer MPH); Leras Afrique, Cotonou, Benin (F G Gankpé MD); CHU Hassan II, Fès, Morocco (F G Gankpé MD); The Task Force for Global Health, Decatur, GA, USA (T Gebre PhD); Ludwig Maximilians University, Munich, Germany (A T Gebremedhin MPH); Division of Human Nutrition (J M Geleijnse PhD), Wageningen University, Wageningen, Netherlands (T Jibat MS); Agence de Médecine Préventive, Paris, France (B D Gessner MD); The Royal Melbourne Hospital, Melbourne, VIC, Australia (K B Gibney FRACP); College of Medicine, University of Hail, Hail, Saudi Arabia (I A Ginawi MD); University Hospital of Dijon, Dijon, France (Prof M Giroud MD); College of Health and Medical Sciences (H S Roba MPH), Haramaya University, Dire Dawa, Ethiopia (M D Gishu MS, A K Tura MPH); Kersa Health and Demographic Surveillance System, Harar, Ethiopia (M D Gishu MS); Heller School for Social Policy and Management (E Glaser PhD), Brandeis University, Waltham, MA, USA (Y A Halasa MS, Prof D S Shepard PhD, E A Undurraga PhD); University of Massachusetts Boston, Boston, MA, USA (Prof P Gona PhD); Instituto de Investigaciones Científicas y Servicios de Alta Tecnología-INDICASAT-AIP, Ciudad del Saber, Panamá (A Goodridge PhD); Department of Health and Social Affairs, Government of the Federated States of Micronesia, Palikir, Federated States of Micronesia (S V Gopalani MPH); University of British Columbia, Vancouver, BC, Canada (C C Gotay PhD, Prof N Kissoon MD, J A Kopec PhD, F Pourmalek PhD); Division of Epidemiology, Center for Public Health Sciences (A Goto PhD), National Cancer Center, Tokyo, Japan (M Inoue MD); Centre for International Health, Dunedin School of Medicine (Prof J A Crump MD), Injury Prevention Research Unit, Department of Preventive and Social Medicine, Dunedin School of Medicine (Prof S Derrett PhD), University of Otago, Wellington, New Zealand (R Grainger PhD); West Virginia Bureau for Public Health, Charleston, WV, USA (R Gupta MD); Eternal Heart Care Centre and Research Institute, Jaipur, India (R Gupta PhD); Department of Anthropology, University of Delhi, Delhi, India (V Gupta PhD); National Institute of Psychiatry Ramon de la Fuente, Mexico City, Mexico (R A Gutiérrez PhD); Department of Global Public Health and Primary Care (A K Knudsen PhD, Prof S E Vollset DrPH), University of Bergen, Bergen, Norway (A D Hailu MPH, Prof O F Norheim PhD); Kiltie Awlaelo Health and Demographic Surveillance System, Mekelle, Ethiopia (G B Hailu MSc); Arabian Gulf University, Manama, Bahrain (Prof R R Hamadeh DPhil); Hamdan Bin Mohammed Smart University, Dubai, United Arab Emirates (S Hamidi PhD); Wayne County Department of Health and Human Services, Detroit, MI, USA (M Hammami MD); University of New Mexico, Albuquerque, NM, USA (A J Handal PhD); School of Medicine and Pharmacology, University of Western Australia, Perth, WA, Australia (Prof G J Hankey MD); Harry Perkins Institute of Medical Research, Nedlands, WA, Australia (Prof G J Hankey MD); Western Australian Neuroscience Research Institute, Nedlands, WA, Australia (Prof G J Hankey MD); School of Public Health, Sun Yat-sen University, Guangzhou, China (Prof Y Hao PhD); Sree Chitra Tirunal Institute for Medical Sciences and Technology, Trivandrum, India (S Harikrishnan DM); Parc Sanitari Sant Joan de Déu - CIBERSAM, Sant Boi de Llobregat (Barcelona), Spain (J M Haro MD); Universitat de Barcelona, Barcelona, Spain (J M Haro MD); International Foundation for Dermatology, London, UK (Prof R J Hay DM); Department of Psychiatry, University Medical Center Groningen (Prof H W Hoek MD), University of Groningen, Groningen, Netherlands (A K Tura MPH); Department of Epidemiology, Mailman School of Public Health (Prof H W Hoek MD), Columbia University, New York, NY, USA (Prof V Skirbekk PhD); Nevada Division of Public and Behavioral Health, Department of Health and Human Services, Carson City, NV, USA (M Horino MPH); Department of Pulmonology, Yokohama City University Graduate School of Medicine, Yokohama, Japan (N Horita MD); Albert Einstein College of Medicine, Bronx, NY, USA (Prof H D Hosgood PhD); Public Health Division, The Pacific Community, Noumea, New Caledonia (D G Hoy PhD); International Relations Division, Ministry of Health, Nay Pyi Taw, Myanmar (A S Htet MPhil); Cambridge Health Alliance, Cambridge, MA, USA (H Huang MD); Aarhus University, Aarhus, Denmark (K M Iburg PhD); National Institute for Health Development, Tallinn, Estonia (K Innos PhD); Graduate School of Medicine (M Inoue MD), School of Public Health (Prof N Kawakami MD), University of Tokyo, Tokyo, Japan (K Shibuya MD); Department of Global and Community Health, George Mason University, Fairfax, VA, USA (K H Jacobsen PhD); Faculty of Medical Sciences, University of Kragujevac, Kragujevac, Serbia (Prof M B Jakovljevic PhD); University of Aberdeen, Aberdeen, UK (M Javanbakht PhD); Department of Surgery, Virginia Commonwealth University, Richmond, VA, USA (S P Jayaraman MD); Postgraduate Institute of Medicine, Colombo, Sri Lanka (A U Jayatilake PhD); Institute of Violence and Injury Prevention, Colombo, Sri Lanka (A U Jayatilake PhD); Graduate School of Public Health, Yonsei University, Seoul, South Korea (Prof S H Jee PhD); Centre for Chronic Disease Control, New Delhi, India (P Jeemon PhD, D Prabhakaran DM); Department of Health Development, Institute of Industrial Ecological Sciences, University of Occupational and Environmental Health, Kitakyushu, Japan (Y Jiang PhD); Department of Ocular Epidemiology and Visual Health, Institute of Ophthalmology Conde de Valencia, Mexico City, Mexico (A Jimenez-Corona PhD); General Directorate of Epidemiology, Ministry of Health, Mexico City, Mexico (A Jimenez-Corona PhD); Department of Ophthalmology, Medical Faculty Mannheim, Ruprecht-Karls-University Heidelberg, Mannheim, Germany (Prof J B Jonas MD); University College Cork, Cork, Ireland (Z Kabir PhD); Society for Education, Action and Research in Community Health, Gadchiroli, India (Y Kalkonde MD); CSIR-Indian Institute of Toxicology Research, Lucknow, India (R Kamal MSc, C N Kesavachandran PhD); Fudan University, Shanghai, China (H Kan MD); Epidemiological and Statistical Methods Research Group, Helmholtz Centre for Infection Research, Braunschweig, Germany (A Karch MD); Hannover-Braunschweig Site, German Center for Infection Research, Braunschweig, Germany (A Karch MD); Quality and Equity Health Care, Kigali, Rwanda (C K Karema MSc); Case Western University Hospitals, Cleveland, OH, USA (C Karimkhani MD); Oklahoma State University, Tulsa, OK, USA (A Kaul MD); Institute of Tropical and Infectious Diseases, Nairobi, Kenya (P N Keiyo PhD); School of Continuing and Distance Education, Nairobi, Kenya (P N Keiyo PhD); Farr Institute (Prof R A Lyons MD), Swansea University, Swansea, UK (Prof A H Kemp PhD); Assuta Hospitals, Assuta Hashalom, Tel Aviv, Israel (Prof A Keren MD); Jordan University

- of Science and Technology, Irbid, Jordan (Prof Y S Khader ScD); Health Services Academy, Islamabad, Pakistan (E A Khan MPH); College of Medicine (Prof Y H Khang MD), Graduate School of Public Health (Prof S Won PhD), Seoul National University, Seoul, South Korea; New York Medical College, Valhalla, NY, USA (S Khera MD, M Tavakkoli MD); Executive Board of the Health Ministers' Council for Cooperation Council States, Riyadh, Saudi Arabia (Prof T A M Khoja FRCP); Ball State University, Muncie, IN, USA (J Khubchandani PhD); Hospital de Clínicas de Porto Alegre, Porto Alegre, Brazil (C Kielsing MD); Korea Health Industry Development Institute, Cheongju-si, South Korea (C Kim PhD); Department of Health Sciences, Northeastern University, Boston, MA, USA (Prof D Kim DrPH); Southern University College, Skudai, Malaysia (Y J Kim PhD); Centre for Disease Burden (A K Knudsen PhD, Prof S E Vollset DrPH), Department of Health Promotion (J C Skogen PhD), Norwegian Institute of Public Health, Oslo, Norway (M Savic PhD, Prof V Skirbekk PhD); Department of Preventive Cardiology, National Cerebral and Cardiovascular Center, Suita, Japan (Y Kokubo PhD); Division of Cardiology (D Kolte MD), Brown University, Providence, RI, USA (Prof S T McGarvey PhD); Center for Community Empowerment, Health Policy and Humanities, NIHRD, Jakarta, Indonesia (S Kosen MD); Sher-i-Kashmir Institute of Medical Sciences, Srinagar, India (Prof P A Koul MD); Research and Development Unit, Parc Sanitari Sant Joan de Deu (CIBERSAM), Barcelona, Spain (A Koyanagi MD); Research Center of Neurology, Moscow, Russia (M Kravchenko PhD, Prof Y Y Varakin MD); Department of Demography and Public Health Research Institute (Prof B Kuate Defo PhD), Department of Social and Preventive Medicine, School of Public Health (Prof B Kuate Defo PhD), University of Montreal, Montreal, QC, Canada; Institute of Public Health, Hacettepe University, Ankara, Turkey (B Kucuk Bicer PhD); University of Cape Coast, Cape Coast, Ghana (A A Kudom PhD); Department of Public Health (S Polinder PhD), Erasmus MC, University Medical Center Rotterdam, Rotterdam, Netherlands (Prof E J Kuipers PhD); Work Organizations, Work Disability Prevention, The Finnish Institute of Occupational Health, Helsinki, Finland (T Lallukka PhD, R Shirir PhD); Department of Public Health, Faculty of Medicine (T Lallukka PhD), University of Helsinki, Helsinki, Finland (T J Meretoja PhD); Institute of Health Policy and Development Studies, National Institutes of Health, Manila, Philippines (Prof H Lam PhD); Johns Hopkins Bloomberg School of Public Health (J O Lam PhD, Prof J B Nachega PhD), Johns Hopkins University, Baltimore, MD, USA (B X Tran PhD); London School of Hygiene & Tropical Medicine, London, UK (S M Langan PhD, Prof M McKee DSc); Servicio de Neurología, Clínica Alemana, Universidad del Desarrollo, Santiago, Chile (P M Lavados MD); College of Optometry, Nova Southeastern University, Fort Lauderdale, FL, USA (J L Leasher OD); State University of New York, Albany, Rensselaer, NY, USA (R Leung PhD); Tuscan Regional Centre for Occupational Injuries and Diseases, Florence, Italy (M Levi PhD); San Francisco VA Medical Center, San Francisco, CA, USA (Y Li PhD); National Office for Maternal and Child Health Surveillance, West China Second University Hospital, Sichuan University, Chengdu, China (Prof J Liang MD); Emory University, Atlanta, GA, USA (Prof Y Liu PhD, Prof M R Phillips MD); Eastern Health Clinical School (B K Lloyd PhD), Monash University, Fitzroy, VIC, Australia; Turning Point, Eastern Health, Melbourne, VIC, Australia (B K Lloyd PhD); Nationwide Children's Hospital, Columbus, OH, USA (W D Lo MD); University of Bari, Bari, Italy (Prof G Logroscino PhD); University of Bristol, Bristol, UK (K J Looker PhD); Aintree University Hospital National Health Service Foundation Trust, Liverpool, UK (Prof R Lunevicius PhD); School of Medicine, University of Liverpool, Liverpool, UK (Prof R Lunevicius PhD); Royal Children's Hospital, Melbourne, VIC, Australia (M T Mackay MBBS, R G Weintraub MBBS); Aswan University Hospital, Aswan Faculty of Medicine, Aswan, Egypt (M Magdy Abd El Razek MBBCh); Social Security Organization Research Institute, Tehran, Iran (M Mahdavi PhD); Institute of Health Policy and Management, Erasmus University Rotterdam, Rotterdam, Netherlands (M Mahdavi PhD); Division of Population and Patient Health, King's College London Dental Institute, London, UK (Prof W Marcenis PhD); Perelman School of Medicine (P A Meaney MD), University of Pennsylvania, Philadelphia, PA, USA (D J Margolis PhD); University Hospital Doctor Peset, University of Valencia, Valencia, Spain (J Martinez-Raga PhD); CEU Cardenal Herrera University, Moncada (Valencia), Spain (J Martinez-Raga PhD); Hospital Pedro Hispano/ULS Matosinhos, Matosinhos, Portugal (J Massano MD); Alaska Native Tribal Health Consortium, Anchorage, AK, USA (B J McMahon MD); Children's Hospital of Philadelphia, Philadelphia, PA, USA (P A Meaney MD); College of Medicine, Howard University, Washington, DC, USA (A Mehari MD); University of Gondar, Gondar, Ethiopia (A B Mekonnen MS, B A Tedla BS); University of West Florida, Pensacola, FL, USA (P Memiah PhD); Saudi Ministry of Health, Riyadh, Saudi Arabia (Prof Z A Memish MD); College of Medicine, Alfaisal University, Riyadh, Saudi Arabia (Prof Z A Memish MD); United Nations Population Fund, Lima, Peru (W Mendoza MD); Department of Neurology, Helsinki University Hospital, Helsinki, Finland (A Meretoja PhD); Helsinki University Hospital, Comprehensive Cancer Center, Breast Surgery Unit, Helsinki, Finland (T J Meretoja PhD); Ifakara Health Institute, Bagamoyo, Tanzania (F A Mhimira MS); Pacific Institute for Research & Evaluation, Calverton, MD, USA (T R Miller PhD); Centre for Population Health, Curtin University, Perth, WA, Australia (T R Miller PhD); University of Ottawa, Ottawa, ON, Canada (E J Mills PhD); Neuroscience Research Center, Baqiyatallah University of Medical Sciences, Tehran, Iran (A Mohammadi PhD); Health Systems and Policy Research Unit, Ahmadu Bello University, Zaria, Nigeria (S Mohammed PhD); Institute for Maternal and Child Health, IRCCS "Burlo Garofolo", Trieste, Italy (L Monasta DSc, M Montico MSc, L Ronfani PhD); Department of Community Medicine, Gastrointestinal and Liver Disease Research Center, Preventive Medicine and Public Health Research Center, Iran University of Medical Sciences, Tehran, Iran (M Moradi-Lakeh MD); International Laboratory for Air Quality and Health (L Morawska PhD), Institute of Health and Biomedical Innovation (R E Norman PhD), Queensland University of Technology, Brisbane, QLD, Australia; Competence Center Mortality-Follow-Up of the German National Cohort (A Werdecker PhD), Federal Institute for Population Research, Wiesbaden, Germany (Prof U O Mueller PhD, R Westerman PhD); Graduate School of Public Health (Prof J B Nachega PhD), Public Health Dynamics Laboratory (A J Paternina Caicedo MD), University of Pittsburgh, Pittsburgh, PA, USA; West Herts Hospitals NHS Trust, Watford, Hertfordshire, UK (M E Murdoch FRCP); Stellenbosch University, Cape Town, South Africa (Prof J B Nachega PhD, Prof S Seedat PhD, Prof C S Wiysonge PhD); Institute of Epidemiology and Medical Biometry, Ulm University, Ulm, Germany (Prof G Nagel PhD, Prof D Rothenbacher MD); International Centre for Diarrhoeal Disease Research, Bangladesh (icddr), Dhaka, Bangladesh (A Naheed PhD); Azienda Ospedaliera Papa Giovanni XXIII, Bergamo, Italy (Prof L Naldi MD, Prof G Remuzzi MD); Suraj Eye Institute, Nagpur, India (V Nangia MD); Ministry of Health and Social Welfare, Dar es Salaam, Tanzania (F N Ngalesoni MSc); Institute for Global Health Innovations, Duy Tan University, Da Nang, Vietnam (Q L Nguyen MD); Institute For Research, Socio-Economic Development and Communication, Yaoundé, Cameroon (P M Nkamedjie Pete MS); Hospital Universitari de Bellvitge, L'Hospitalet, Spain (J M Nolla PhD); Federal University of Pelotas, Pelotas, Brazil (Prof B P Nunes PhD); Centre for Health Research, Western Sydney University, Sydney, NSW, Australia (F A Ogbo MPH); Department of Preventive Medicine, School of Medicine, Kyung Hee University, Seoul, South Korea (Prof I Oh PhD); Teikyo University School of Medicine, Tokyo, Japan (Prof T Ohkubo MD); Universidad Autonoma de Chile, Talca, Chile (Prof P R Olivares PhD); Center for Healthy Start Initiative, Lagos, Nigeria (B O Olusanya PhD, J O Olusanya MBA); IIS-Fundacion Jimenez Diaz-UAM, Madrid, Spain (Prof A Ortiz PhD); YBank, Cambridge, MA, USA (M Osman MD); St Luke's International University, Tokyo, Japan (E Ota PhD); JSS Medical College, JSS University, Mysore, India (Prof Mahesh PA DNB); Department of Medical Humanities and Social Medicine, College of Medicine, Kosin University, Busan, South Korea (E Park PhD); Universidade Federal de Minas Gerais, Belo Horizonte, Brazil (Prof V M D A Passos PhD); Universidad de Cartagena, Cartagena, Colombia (A J Paternina Caicedo MD); Department of Community Health Sciences (Prof S B Patten PhD), University of Calgary, Calgary, AB, Canada (Prof M Tonelli MD); REQUIMTE/LAQV, Laboratório de

Farmacognosia, Departamento de Química, Faculdade de Farmácia, Universidade do Porto, Porto, Portugal (Prof D M Pereira PhD); National Institute of Respiratory Diseases, Mexico City, Mexico (Prof R Perez-Padilla MD); Flinders University, Adelaide, SA, Australia (Prof K Pesudovs PhD); University of the Witwatersrand, Johannesburg, South Africa (Prof M Petzold PhD); Shanghai Jiao Tong University School of Medicine, Shanghai, China (Prof M R Phillips MD); Durban University of Technology, Durban, South Africa (J D Pillay PhD); Exposure Assessment and Environmental Health Indicators, German Environment Agency, Berlin, Germany (D Plass DrPH); Department of Anesthesiology (A S Terkawi MD), University of Virginia, Charlottesville, VA, USA (J A Platts-Mills MD); University of Newcastle, Callaghan, NSW, Australia (Prof C D Pond PhD); The Fred Hollows Foundation, Sydney, NSW, Australia (N M Prasad DO); Centre for Eye Research Australia, Melbourne, VIC, Australia (N M Prasad DO); Department of Community Medicine, School of Medicine, Alborz University of Medical Sciences, Karaj, Iran (M Qorbani PhD); A T Still University, Kirksville, MO, USA (A Radfar MD); Contech School of Public Health, Lahore, Pakistan (A Rafay MS); Research and Evaluation Division, BRAC, Dhaka, Bangladesh (M Rahman PhD); Hamad Medical Corporation, Doha, Qatar (S U Rahman FCPS); Society for Health and Demographic Surveillance, Suri, India (R K Rai MPH); ERAWEB Program, University for Health Sciences, Medical Informatics and Technology, Hall in Tirol, Austria (S Rajsic MD); Walden University, Minneapolis, MN, USA (Prof A H Refaat PhD); Suez Canal University, Ismailia, Egypt (Prof A H Refaat PhD); Department of Biomedical and Clinical Sciences L Sacco, University of Milan, Milan, Italy (Prof G Remuzzi MD); Hospital das Clinicas da Universidade Federal de Minas Gerais, Belo Horizonte, Brazil (Prof A L Ribeiro MD); (ISGlobal) Instituto de Salud Global de Barcelona, Barcelona, Spain (D Rojas-Rueda PhD); Golestan Research Center of Gastroenterology and Hepatology, Golestan University of Medical Sciences, Gorgan, Iran (G Roshandel PhD); All India Institute of Medical Sciences, New Delhi, India (A Roy DM, R Sagar MD, M Satpathy PhD, Prof N Tandon PhD); Ballarat Health Service, Ballarat, VIC, Australia (R Sahathevan PhD); Universiti Kebangsaan Malaysia Medical Centre, Kuala Lumpur, Malaysia (R Sahathevan PhD); Marshall University J Edwards School of Medicine, Huntington, WV, USA (J R Sanabria MD); Case Western Reserve University, Cleveland, OH, USA (J R Sanabria MD); IIS-Fundacion Jimenez Diaz, Madrid, Spain (M D Sanchez-Niño PhD); Universidad Ciencias Aplicadas y Ambientales, Bogotá, Colombia (R Sarmiento-Suarez MPH); University of KwaZulu-Natal, Durban, South Africa (Prof B Sartorius PhD); Marshall University, Huntington, WV, USA (M Sawhney PhD); Swiss Research Institute of Public Health and Addiction (M P Schaub PhD), University of Zurich, Zurich, Switzerland (H G Yebo MS); Federal University of Santa Catarina, Florianópolis, Brazil (J C Schneider PhD, D A S Silva PhD); German Cancer Research Center, Heidelberg, Germany (B Schöttker MPH); Institute of Health Care and Social Sciences, FOM University, Essen, Germany (B Schöttker MPH); University of Alabama at Birmingham, Birmingham, AL, USA (D C Schwebel PhD, J A Singh MD); Department of Public Health, An-Najah University, Nablus, Palestine (A Shaheen PhD); Independent Consultant, Karachi, Pakistan (M A Shaikh MD); Indian Institute of Technology Ropar, Rupnagar, India (R Sharma MA); ICMR National Institute of Epidemiology, Chennai, India (U Sharma MPH); Research Institute at Nationwide Children's Hospital, Columbus, OH, USA (J Shen PhD); Department of Public Health Science, Graduate School (Prof M Shin PhD), Department of Preventive Medicine, College of Medicine (S Yoon PhD), Korea University, Seoul, South Korea; Faculty of Health and Life Sciences, Northumbria University, Newcastle upon Tyne, UK (I Shiue PhD); Reykjavik University, Reykjavik, Iceland (I D Sigfusdottir PhD); Brasília University, Brasília, Brazil (D G A Silveira MD); Department of Medicine, Institute of Medical Sciences, Banaras Hindu University, Varanasi, India (O P Singh PhD); Institute for Human Development, New Delhi, India (P K Singh PhD); Alcohol and Drug Research Western Norway (J C Skogen PhD), Stavanger University Hospital, Stavanger, Norway (K Søreide PhD); Faculty of Health Sciences, Hatter Institute for Cardiovascular Research in Africa (Prof K Sliwa PhD), Department of Psychiatry (Prof D J Stein PhD), University of Cape Town, Cape Town, South Africa (D A Watkins MD); Instituto de Investigación Hospital Universitario de la Princesa, Universidad Autónoma de Madrid, Cátedra UAM-Linde, Palma de Mallorca, Spain (Prof J B Soriano PhD); Department of Clinical Neurological Sciences, Western University, London, ON, Canada (L A Sposato MD); Department of Community Medicine, International Medical University, Kuala Lumpur, Malaysia (C T Sreeramareddy MD); Attikon University Hospital, Athens, Greece (V Stathopoulou PhD); University of East Anglia, Norwich, UK (Prof N Steel PhD); South African Medical Research Council Unit on Anxiety & Stress Disorders, Cape Town, South Africa (Prof D J Stein PhD); Department of Neuroscience, Norwegian University of Science and Technology, Trondheim, Norway (Prof T J Steiner PhD, Prof L J Stovner PhD); Department of Dermatology, University Hospital Muenster, Muenster, NRW, Germany (S Steinke DrMed); Norwegian Advisory Unit on Headache, St Olavs Hospital, Trondheim, Norway (Prof L J Stovner PhD); Alexandra General Hospital of Athens, Athens, Greece (K Stroumpoulis PhD); Centre Hospitalier Public du Cotentin, Cherbourg, France (K Stroumpoulis PhD); Muhimbili University of Health and Allied Sciences, Dar es Salaam, Tanzania (B F Sunguya PhD); Indian Council of Medical Research, New Delhi, India (S Swaminathan MD); Departments of Criminology, Law & Society, Sociology, and Public Health, University of California, Irvine, Irvine, CA, USA (Prof B L Sykes PhD); Department of Medicine, University of Valencia, INCLIVA Health Research Institute and CIBERSAM, Valencia, Spain (Prof R Tabarés-Seisdedos PhD); WSH Institute, Ministry of Manpower, Singapore, Singapore (J S Takala DSc); Tampere University of Technology, Tampere, Finland (J S Takala DSc); Chaim Sheba Medical Center, Tel Hashomer, Israel (Prof D Tanne MD); Tel Aviv University, Tel Aviv, Israel (Prof D Tanne MD); James Cook University, Cairns, QLD, Australia (B A Tedla BS); Outcomes Research Consortium (A S Terkawi MD), Cleveland Clinic, Cleveland, OH, USA (Prof E M Tuzcu MD); Department of Anesthesiology, King Fahad Medical City, Riyadh, Saudi Arabia (A S Terkawi MD); Adaptive Knowledge Management, Victoria, BC, Canada (A J Thomson PhD); WorldFish, Penang, Malaysia (A L Thorne-Lyman ScD); Nelson Institute of Environmental Medicine, School of Medicine (Prof G D Thurston ScD), New York University, Tuxedo, NY, USA; National Center for Child Health and Development, Tokyo, Japan (R Tobe-Gai PhD); Institute of Public Health, Faculty of Health Sciences, Jagiellonian University Medical College, Kraków, Poland (R Topor-Madry PhD); Faculty of Health Sciences, Wrocław Medical University, Wrocław, Poland (R Topor-Madry PhD); Aristotle University of Thessaloniki, Thessaloniki, Greece (Prof F Topouzis PhD); Hanoi Medical University, Hanoi, Vietnam (B X Tran PhD); Department of Neurology, Rigshospitalet, University of Copenhagen, Denmark (T Truelsen DMSc); Department of Population Sciences and Development, Faculty of Economics and Management, University of Kinshasa, Kinshasa, Democratic Republic of the Congo (Z Tsala Dimbuene PhD); African Population and Health Research Center, Nairobi, Kenya (Z Tsala Dimbuene PhD); Department of Medicine, University of Crete, Heraklion, Greece (Prof M Tsilimbaris PhD); Parc Sanitari Sant Joan de Déu, Fundació Sant Joan de Déu, Universitat de Barcelona, CIBERSAM, Barcelona, Spain (S Tyrovolas PhD); Department of Internal Medicine, Federal Teaching Hospital, Abakaliki, Nigeria (K N Ukwaja MD); Ebonyi State University, Abakaliki, Nigeria (C J Ukeke PhD); Warwick Medical School, University of Warwick, Coventry, UK (O A Uthman PhD); National Institute for Public Health and the Environment, Bilthoven, Netherlands (C H van Gool PhD); UKK Institute for Health Promotion Research, Tampere, Finland (Prof T Vasankari PhD); Raffles Neuroscience Centre, Raffles Hospital, Singapore, Singapore (N Venketasubramanian FRCP); University of Bologna, Bologna, Italy (Prof F S Violante MD); Federal Research Institute for Health Organization and Informatics, Moscow, Russia (S K Vladimirov PhD); National Research University Higher School of Economics, Moscow, Russia (Prof V V Vlassov MD); National Institute for Occupational Safety and Health, Washington, DC, USA (G R Wagner MD); Uniformed Services University of Health Sciences, Bethesda, MD, USA (Prof S G Waller MD); McGill University, Montreal, QC, Canada (S Weichenthal PhD); Department of Research, Cancer Registry of Norway, Institute of Population-Based Cancer Research, Oslo, Norway (E Weiderpass PhD); Department of Community Medicine,

Faculty of Health Sciences, University of Tromsø, The Arctic University of Norway, Tromsø, Norway (E Weiderpass PhD); Genetic Epidemiology Group, Folkhälsan Research Center, Helsinki, Finland (E Weiderpass PhD); German National Cohort Consortium, Heidelberg, Germany (R Westerman PhD); Department of Infectious Disease Epidemiology and Modelling (R A White PhD), Norwegian Institute of Public Health, Oslo, Norway (M Savic PhD, Prof V Skirbekk PhD); Centre of Evidence-based Dermatology, University of Nottingham, Nottingham, UK (Prof H C Williams DSc); South African Medical Research Council, Cape Town, South Africa (Prof C S Wiysonge PhD); National Institute for Health Research Comprehensive Biomedical Research Centre, Guy's & St Thomas' NHS Foundation Trust and King's College London, London, UK (Prof C D Wolfe MD); St Paul's Hospital, Millennium Medical College, Addis Ababa, Ethiopia (M Wubshet PhD); St John's Medical College and Research Institute, Bangalore, India (Prof D Xavier MD); Department of Neurology, Jinling Hospital, Nanjing University School of Medicine, Nanjing, China (Prof G Xu PhD); Global Health Research Center, Duke Kunshan University, Kunshan, China (Prof L L Yan PhD); Department of Preventive Medicine, Northwestern University, Chicago, IL, USA (Y Yano MD); Social Work and Social Administration Department and The Hong Kong Jockey Club Centre for Suicide Research and Prevention, University of Hong Kong, Hong Kong, China (Prof P Yip PhD); Department of Biostatistics, School of Public Health, Kyoto University, Kyoto, Japan (N Yonemoto MPH); Jackson State University, Jackson, MS, USA (Prof M Z Younis DrPH); Department of Epidemiology and Biostatistics, School of Public Health (Prof C Yu PhD), Global Health Institute (Prof C Yu PhD), Wuhan University, Wuhan, China; University Hospital, Setif, Algeria (Prof Z Zaidi PhD); Faculty of Medicine, Mansoura University, Mansoura, Egypt (Prof M E Zaki PhD); Leibniz Institute for Prevention Research and Epidemiology, Bremen, Germany (Prof H Zeeb PhD); Red Cross War Memorial Children's Hospital, Cape Town, South Africa (L J Zuhlke PhD).

Contributors

Christopher J L Murray and Theo Vos prepared the first draft. Alan D Lopez and Christopher J L Murray conceived the study and provided overall guidance. All other authors provided data, developed models, reviewed results, initiated modelling infrastructure, and/or reviewed and contributed to the report.

Declaration of interests

Bruce Bartholow Duncan and Maria Inês Schmidt have received additional funding from the Brazilian Ministry of Health (Process No 25000192049/2014-14). Itamar S Santos reports grants from FAPESP (Brazilian public agency), outside the submitted work. Carl Abelardo T Antonio reports grants, personal fees and non-financial support from Johnson & Johnson (Philippines), Inc, outside the submitted work. Cyrus Cooper reports other from Alliance for Better Bone Health, other from Amgen, other from Eli Lilly, other from GSK, other from Medtronic, other from Merck, other from Novartis, other from Pfizer, other from Roche, other from Servier, outside the submitted work. Walter Mendoza is currently employed by the Peru Country Office of the United Nations Population Fund, an institution which does not necessarily endorse this study. Donald S Shepard would like to acknowledge grant support from Sanofi Pasteur. Rafael Tabarés-Seisdedos and Ferrán Catalá-López are supported in part by grant PROMETEOII/2015/021 from Generalitat Valenciana, and Rafael Tabarés-Seisdedos is supported by the national grant PI14/00894 from ISCIII-FEDER. Walter Mendoza is currently employed by the Peru Country Office of the United Nations Population Fund, an institution which does not necessarily endorse this study. Veena J Iyer has received a Public Health Research Initiative Fellowship (2014-17) from the Department of Science and Technology (DST) for a project titled "Relationship between Enteric Fever incidence and Climate in the city of Ahmedabad, 1990–2014". Pablo M Lavados reports grants, personal fees and non-financial support from BAYER, non-financial support from Boehringer Ingelheim, grants and personal fees from AstraZeneca, grants from CONICYT, and grants from The George Institute for Global Health, outside the submitted work. Noela M Prasad reports and is an employee of an international NGO that raises funds from the Australian Public, and holds a honorary position at CERA, which receives Operational Infrastructure Support from the Victorian Government.

Bradford D Gessner reports grants from Crucell, GSK, Hilleman Labs, Novartis, Pfizer, Merck, and Sanofi Pasteur, outside the submitted work. Ai Koyanagi's work is supported by the Miguel Servet contract financed by the CP13/00150 and PI15/00862 projects, integrated into the National R + D + I and funded by the ISCIII - General Branch Evaluation and Promotion of Health Research - and the European Regional Development Fund (ERDF-FEDER). Dorairaj Prabhakaran reports grants from the Bill & Melinda Gates Foundation. Matthias Endres reports that The Center for Stroke Research Berlin has received institutional funding from the German Ministry for Research and Education (BMBF). Katharine J Looker has received funding from the World Health Organization for the HSV-2 seroprevalence review which informs this work; during the study, KJL also received separate funding from the World Health Organization, USAID/PATH, Sexual Health 24 and the National Institute for Health Research (NIHR) Health Protection Research Unit (HPRU) in Evaluation of Interventions at the University of Bristol; these funders had no role in the writing of the manuscript nor the decision to submit it for publication, and the views expressed in this Article do not necessarily represent the views, decisions or policies of the World Health Organization, the NHS, the NIHR, the Department of Health or Public Health England. Donal Bisanzio is supported by Bill and Melinda Gates Foundation (#OPP1068048). Thomas Fürst has received financial support from the Swiss National Science Foundation (SNSF; project no P300P3-154634). Rodrigo Sarmiento-Suarez receives institutional support from Universidad de Ciencias Aplicadas y Ambientales, UDCA, Bogotá Colombia. Ronan A Lyons is supported by two grants: The Farr Institute of Health Informatics Research: Arthritis Research UK, the British Heart Foundation, Cancer Research UK, the Economic and Social Research Council, the Engineering and Physical Sciences Research Council, the Medical Research Council, the National Institute of Health Research, the National Institute for Social Care and Health Research (Welsh Assembly Government), the Chief Scientist Office (Scottish Government Health Directorates), and The Wellcome Trust. Grant No MR/K006525/1, and the National Centre for Population Health and Wellbeing Research. Health and Care Research Wales. Stefanos Tyrovolas's work is supported by the Foundation for Education and European Culture (IPEP), the Sara Borrell postdoctoral programme (reference no CD15/00019 from the Instituto de Salud Carlos III (ISCIII - Spain) and the Fondos Europeo de Desarrollo Regional (FEDER). Manami Inoue is the beneficiary of a financial contribution from the AXA Research fund as chair holder of the AXA Department of Health and Human Security, Graduate School of Medicine, The University of Tokyo from Nov 1, 2012; the AXA Research Fund has no role in this work. Sinead M Langan holds an NIHR Clinician Scientist Fellowship (NIHR/CS/010/014); the views expressed in this publication are those of the authors and not necessarily those of the NHS, the NIHR, or the UK Department of Health. Scott Weichenthal acknowledges financial support from the Cancer Research Society of Canada. Yogeshwar Kalkonde is a Wellcome Trust/ DBT India Alliance Intermediate Fellow in Public Health. John J McGrath received a NHMRC John Cade Fellowship (APP1056929). Sarah Derrett reports grants, personal fees, non-financial support and other from EuroQol Research Foundation, outside the submitted work. Dan J Stein reports personal fees from Lundbeck, personal fees from Novartis, personal fees from AMBRF, grants from NRGF, personal fees from Biocodex, personal fees from Sevier, grants from MRC, personal fees from SUN, and personal fees from CIPLA, outside the submitted work. Tea Lallukka reports funding from The Academy of Finland, grant #287488. Charles D A Wolfe's research was funded and supported by the National Institute for Health Research (NIHR) Biomedical Research Centre based at Guy's and St Thomas' NHS Foundation Trust and King's College London. The views expressed are those of the author(s) and not necessarily those of the NHS, the NIHR, or the Department of Health. The other authors declare no competing interests.

Acknowledgments

We would like to thank the countless individuals who have contributed to the Global Burden of Disease Study 2015 in various capacities. This paper uses data from SHARE Waves 1, 2, 3 (SHARELIFE), 4, and 5 (DOIs: 10.6103/SHARE.w1.500, 10.6103/SHARE.w2.500, 10.6103/SHARE.w3.500, 10.6103/SHARE.w4.500, 10.6103/SHARE.w5.500), see Börsch-Supan et al (2013) for methodological details. The SHARE data collection has been

primarily funded by the European Commission through FP5 (QLK6-CT-2001-00360), FP6 (SHARE-13: RII-CT-2006-062193, COMPARE: CIT5-CT-2005-028857, SHARELIFE: CIT4-CT-2006-028812), and FP7 (SHARE-PREP: N°211909, SHARE-LEAP: N°227822, SHARE M4: N°261982). Additional funding from the German Ministry of Education and Research, the US National Institute on Aging (U01_AG09740-13S2, P01_AG005842, P01_AG08291, P30_AG12815, R21_AG025169, Y1-AG-4553-01, IAG_BSR06-11, OGHA_04-064) and from various national funding sources is gratefully acknowledged (see www.share-project.org). The data reported here have been supplied by the United States Renal Data System (USRDS). The interpretation and reporting of these data are the responsibility of the author(s) and in no way should be seen as an official policy or interpretation of the US Government. The Palestinian Central Bureau of Statistics granted the researchers access to relevant data in accordance with license no. SLN2014-3-170, after subjecting data to processing aiming to preserve the confidentiality of individual data in accordance with the General Statistics Law—2000. The researchers are solely responsible for the conclusions and inferences drawn upon available data. This study has been realised using the data collected by the Swiss Household Panel (SHP), which is based at the Swiss Centre of Expertise in the Social Sciences FORS. The project is financed by the Swiss National Science Foundation. The following individuals would like to acknowledge various forms of institutional support: Amanda G Thrift is supported by a fellowship from the National Health and Medical Research Council (GNT1042600). Panniyammakal Jeemon is supported by the Wellcome Trust-DBT India Alliance, Clinical and Public Health, Intermediate Fellowship (2015–20). Amador Goodridge would like to acknowledge funding from me from Sistema Nacional de Investigadores de Panamá-SNI. José das Neves was supported in his contribution to this work by a Fellowship from Fundação para a Ciência e a Tecnologia, Portugal (SFRH/BPD/92934/2013). Boris Bikbov, Monica Cortinovis, Giuseppe Remuzzi, and Norberto Perico would like to acknowledge that their contribution to this paper has been on behalf of the International Society of Nephrology (ISN) as a follow-up of the activities of the GBD 2010 Genitourinary Diseases Expert Group. Lijing L Yan is supported by the National Natural Sciences Foundation of China grants (71233001 and 71490732). Miriam Levi would like to acknowledge the institutional support received from CeRIMP, Regional Centre for Occupational Diseases and Injuries, Tuscany Region, Florence, Italy. Simon I Hay is funded by a Senior Research Fellowship from the Wellcome Trust (#095066), and grants from the Bill & Melinda Gates Foundation (OPP1119467, OPP1093011, OPP1106023 and OPP1132415). No individuals acknowledged received additional compensation for their efforts.

References

- Atun R, de Jongh T, Secci F, Ohiri K, Adeyi O. Integration of targeted health interventions into health systems: a conceptual framework for analysis. *Health Policy Plan* 2010; **25**: 104–11.
- Park J-H, Eum J-H, Bold B, Cheong H-K. Burden of disease due to dementia in the elderly population of Korea: present and future. *BMC Public Health* 2013; **13**: 293.
- George-Carey R, Adeloye D, Chan KY, et al. An estimate of the prevalence of dementia in Africa: a systematic analysis. *J Glob Health* 2012; **2**: 020401.
- Global Burden of Disease 2013 Collaborators. Global, regional, and national incidence, prevalence, and years lived with disability for 301 acute and chronic diseases and injuries in 188 countries, 1990–2013: a systematic analysis for the Global Burden of Disease Study 2013. *Lancet* 2015; **386**: 743–800.
- Prince MJ, Wu F, Guo Y, et al. The burden of disease in older people and implications for health policy and practice. *Lancet* 2015; **385**: 549–62.
- Hallett TB, Zaba B, Stover J, et al. Embracing different approaches to estimating HIV incidence, prevalence and mortality. *AIDS* 2014; **28**: S523–32.
- Supervie V, Archibald CP, Costagliola D, et al. GBD 2013 and HIV incidence in high income countries. *Lancet* 2015; **385**: 1177.
- Cohen J, Vincent J-L, Adhikari NKJ, et al. Sepsis: a roadmap for future research. *Lancet Infect Dis* 2015; **15**: 581–614.
- Marshall JC. Understanding the global burden of pediatric sepsis. *Am J Respir Crit Care Med* 2015; **191**: 1096–98.
- Anderson BO, Flanigan J. Novel methods for measuring global cancer burden: implications for global cancer control. *JAMA Oncol* 2015; **1**: 425–27.
- Ton TGN, Mackenzie C, Molyneux DH. The burden of mental health in lymphatic filariasis. *Infect Dis Poverty* 2015; **4**: 34.
- Wagner RG, Ibinda F, Tollman S, Lindholm L, Newton CR, Bertram MY. Differing methods and definitions influence DALY estimates: using population-based data to calculate the burden of convulsive epilepsy in rural South Africa. *PLoS One* 2015; **10**: e0145300.
- Alzheimer's Disease International. World Alzheimer Report, 2015. <https://www.alz.co.uk/research/WorldAlzheimerReport2015.pdf> (accessed May 26, 2016).
- Hausman DM. Health, well-being, and measuring the burden of disease. *Popul Health Metr* 2012; **10**: 13.
- Nord E. Uncertainties about disability weights for the Global Burden of Disease study. *Lancet Glob Health* 2015; **3**: e661–62.
- GBD Mortality and Causes of Death Collaborators. Global, regional, and national life expectancy, all-cause and cause-specific mortality for 249 causes of death, 1980–2015: a systematic analysis for the Global Burden of Disease Study 2015. *Lancet* 2015; **388**: 1459–544.
- Stevens GA, Alkema L, Black RE, et al. Guidelines for Accurate and Transparent Health Estimates Reporting: The GATHER statement. *Lancet* 2016; published online June 28. [http://dx.doi.org/10.1016/S0140-6736\(16\)30388-9](http://dx.doi.org/10.1016/S0140-6736(16)30388-9).
- Kiyono P. An integrative meta-regression framework for descriptive epidemiology, 1 edn. Seattle: University of Washington Press, 2015.
- Clark DV, Kibuuka H, Millard M, et al. Long-term sequelae after Ebola virus disease in Bundibugyo, Uganda: a retrospective cohort study. *Lancet Infect Dis* 2015; **15**: 905–12.
- Qureshi AI, Chughtai M, Loua TO, et al. Study of Ebola virus disease survivors in Guinea. *Clin Infect Dis* 2015; **61**: 1035–42.
- Rowe AK, Bertolli J, Khan AS, et al. Clinical, virologic, and immunologic follow-up of convalescent Ebola hemorrhagic fever patients and their household contacts, Kikwit, Democratic Republic of the Congo. *J Infect Dis* 1999; **179**: S28–35.
- Bwaka MA, Bonnet M-J, Calain P, et al. Ebola Hemorrhagic Fever in Kikwit, Democratic Republic of the Congo: clinical observations in 103 patients. *J Infect Dis* 1999; **179**: S1–S7.
- PRO-ACT. <https://nctu.partners.org/ProACT/Data/Index?Length=0&LongLength=0&Rank=1&SyncRoot=System.Type%5B%5D&IsReadOnly=False&IsFixedSize=True&IsSynchronized=False> (accessed May 26, 2016).
- Vos T, Flaxman AD, Naghavi M, et al. Years lived with disability (YLDs) for 1160 sequelae of 289 diseases and injuries 1990–2010: a systematic analysis for the Global Burden of Disease Study 2010. *Lancet* 2012; **380**: 2163–96.
- Global, regional, and national age-sex specific all-cause and cause-specific mortality for 240 causes of death, 1990–2013: a systematic analysis for the Global Burden of Disease Study 2013. *Lancet* 2015; **385**: 117–71.
- Murray CJL, Barber RM, Foreman KJ, et al. Global, regional, and national disability-adjusted life years (DALYs) for 306 diseases and injuries and healthy life expectancy (HALE) for 188 countries, 1990–2013: quantifying the epidemiological transition. *Lancet* 2015; **386**: 2145–91.
- United Nations Development Programme. Human development report 2015. http://hdr.undp.org/sites/default/files/2015_human_development_report.pdf (accessed May 26, 2016).
- Goldman DP, Cutler D, Rowe JW, et al. Substantial health and economic returns from delayed aging may warrant a new focus for medical research. *Health Aff (Millwood)* 2013; **32**: 1698–705.
- Dufouil C, Pereira E, Chêne G, et al. Older age at retirement is associated with decreased risk of dementia. *Eur J Epidemiol* 2014; **29**: 353–61.
- Belbase A, Sanzenbacher G, Gillis CM. Does age-related decline in ability correspond with retirement age? Rochester, NY: Social Science Research Network, 2015 <http://papers.ssrn.com/abstract=2665830> (accessed June 24, 2016).
- van Rijn RM, Robroek SJW, Brouwer S, Burdorf A. Influence of poor health on exit from paid employment: a systematic review. *Occup Environ Med* 2014; **71**: 295–301.
- Lahelma E, Pietiläinen O, Rahkonen O, Lallukka T. Common mental disorders and cause-specific disability retirement. *Occup Environ Med* 2015; **72**: 181–87.

- 33 Begg S, Vos T, Goss J, Mann N. An alternative approach to projecting health expenditure in Australia. *Aust Health Rev* 2008; 32: 148–55.
- 34 Desveaux L, Beauchamp M, Goldstein R, Brooks D. Community-based exercise programs as a strategy to optimize function in chronic disease: a systematic review. *Med Care* 2014; 52: 216–26.
- 35 Jahanbin I, Hoseini Moghadam M, Nazarinia MA, Ghodsbin F, Bagheri Z, Ashraf AR. The effect of conditioning exercise on the health status and pain in patients with rheumatoid arthritis: a randomized controlled clinical trial. *Int J Community Based Nurs Midwifery* 2014; 2: 169–76.
- 36 Gay C, Chabaud A, Guille E, Coudeyre E. Educating patients about the benefits of physical activity and exercise for their hip and knee osteoarthritis. Systematic literature review. *Ann Phys Rehabil Med* 2016; 59: 174–83.
- 37 McAlindon TE, Bannuru RR, Sullivan MC, et al. OARSI guidelines for the non-surgical management of knee osteoarthritis. *Osteoarthritis Cartilage* 2014; 22: 363–88.
- 38 Heimans L, Akdemir G, Boer KVCW, et al. Two-year results of disease activity score (DAS)-remission-steered treatment strategies aiming at drug-free remission in early arthritis patients (the IMPROVED-study). *Arthritis Res Ther* 2016; 18: 23.
- 39 Kleinman A, Estrin GL, Usmani S, et al. Time for mental health to come out of the shadows. *Lancet* 2016; 387: 2274–75.
- 40 Izutsu T, Tsutsumi A, Minas H, Thornicroft G, Patel V, Ito A. Mental health and wellbeing in the Sustainable Development Goals. *Lancet Psychiatry* 2015; 2: 1052–54.
- 41 Patel V, Chisholm D, Parikh R, et al. Addressing the burden of mental, neurological, and substance use disorders: key messages from Disease Control Priorities, 3rd edn. *Lancet* 2016; 387: 1672–85.
- 42 Centers for Disease Control and Prevention. Drug-poisoning deaths involving heroin: United States, 2000–2013. <http://www.cdc.gov/nchs/products/databriefs/db190.htm> (accessed June 24, 2016).
- 43 Centers for Disease Control and Prevention. Morbidity and Mortality Weekly Report (MMWR). <https://www.cdc.gov/mmwr/preview/mmwrhtml/mm6401a10.htm> (accessed June 24, 2016).
- 44 Centers for Disease Control and Prevention. Opioid overdose prevention programs providing naloxone to laypersons—United States, 2014. http://origin.glb.cdc.gov/mmwr/preview/mmwrhtml/mm6423a2.htm?s_cid=mm6423a2_w (accessed June 24, 2016).
- 45 Mathers BM, Degenhardt L, Ali H, et al. HIV prevention, treatment, and care services for people who inject drugs: a systematic review of global, regional, and national coverage. *Lancet* 2010; 375: 1014–28.
- 46 GBD 2015 Risk Factors Collaborators. Global, regional, and national comparative risk assessment of 79 behavioural, environmental and occupational, and metabolic risks or clusters of risks, 1990–2015: a systematic analysis for the Global Burden of Disease Study 2015. *Lancet* 2016; 388: 1659–724.
- 47 Association AD. Economic costs of diabetes in the US in 2012. *Diabetes Care* 2013; 36: 1033–46.
- 48 NCD Risk Factor Collaboration. Worldwide trends in diabetes since 1980: a pooled analysis of 751 population-based studies with 4·4 million participants. *Lancet* 2016; 387: 1513–30.
- 49 International Diabetes Federation. IDF diabetes atlas. <http://www.diabetesatlas.org/> (accessed June 25, 2016).
- 50 Langa KM. Is the risk of Alzheimer's disease and dementia declining? *Alzheimers Res Ther* 2015; 7: 34.
- 51 Matthews FE, Arthur A, Barnes LE, et al. A two-decade comparison of prevalence of dementia in individuals aged 65 years and older from three geographical areas of England: results of the Cognitive Function and Ageing Study I and II. *Lancet* 2013; 382: 1405–12.
- 52 Rocca WA, Petersen RC, Knopman DS, et al. Trends in the incidence and prevalence of Alzheimer's disease, dementia, and cognitive impairment in the United States. *Alzheimers Dement* 2011; 7: 80–93.
- 53 Brookmeyer R, Evans DA, Hebert L, et al. National estimates of the prevalence of Alzheimer's disease in the United States. *Alzheimers Dement* 2011; 7: 61–73.
- 54 Helsper CW, Hellinga HL, van Essen GA, et al. Real-life costs of hepatitis C treatment. *Neth J Med* 2012; 70: 145–53.
- 55 Bhatt S, Weiss DJ, Cameron E, et al. The effect of malaria control on *Plasmodium falciparum* in Africa between 2000 and 2015. *Nature* 2015; 526: 207–11.
- 56 Cameron E, Battle KE, Bhatt S, et al. Defining the relationship between infection prevalence and clinical incidence of *Plasmodium falciparum* malaria. *Nat Commun* 2015; 6: 8170.
- 57 Bhatt S, Weiss DJ, Mappin B, et al. Coverage and system efficiencies of insecticide-treated nets in Africa from 2000 to 2017. *eLife* 2015; 4: e09672.
- 58 Aspiration to action. What will it take to end malaria? <http://endmalaria2040.org/assets/Aspiration-to-Action.pdf> (accessed May 26, 2016).
- 59 WHO. World Malaria Report 2015. <http://www.who.int/malaria/publications/world-malaria-report-2015/report/en/> (accessed Aug 5, 2016).
- 60 Globocan 2012. <http://globocan.iarc.fr/Default.aspx> (accessed June 24, 2016).
- 61 Pizzarello L, Abiose A, Ffytche T, et al. VISION 2020: The Right to Sight: a global initiative to eliminate avoidable blindness. *Arch Ophthalmol* 1960 2004; 122: 615–20.
- 62 Baltussen R, Smith A. Cost effectiveness of strategies to combat vision and hearing loss in sub-Saharan Africa and South East Asia: mathematical modelling study. *BMJ* 2012; 344: e615.
- 63 Currie J, Grenfell B, Farrar J. Beyond Ebola. *Science* 2016; 351: 815–16.
- 64 Baize S, Pannetier D, Oestereich L, et al. Emergence of Zaire Ebola virus disease in Guinea. *N Engl J Med* 2014; 371: 1418–25.
- 65 Weaver SC. Arrival of Chikungunya virus in the new world: prospects for spread and impact on public health. *PLoS Negl Trop Dis* 2014; 8: e2921.
- 66 Fauci AS, Morens DM. Zika Virus in the Americas—yet another arbovirus threat. *N Engl J Med* 2016; 374: 601–04.
- 67 Microcephaly in Infants, Pernambuco State, Brazil, 2015. *Emerg Infect Dis* 2016; 22: 1090–93.
- 68 Rasmussen SA, Jamieson DJ, Honein MA, Petersen LR. Zika Virus and birth defects—reviewing the evidence for causality. *N Engl J Med* 2016; 374: 1981–87.
- 69 Jensen ET, Cook SF, Allen JK, et al. Enrollment factors and bias of disease prevalence estimates in administrative claims data. *Ann Epidemiol* 2015; 25: 519–25.
- 70 Kottke TE, Baechler CJ, Parker ED. Accuracy of heart disease prevalence estimated from claims data compared with an electronic health record. *Prev Chronic Dis* 2012; 9: E141.
- 71 Salomon JA, Nordhagen S, Oza S, Murray CJL. Are Americans feeling less healthy? The puzzle of trends in self-rated health. *Am J Epidemiol* 2009; 170: 343–51.
- 72 Salomon JA, Tandon A, Murray CJL. Comparability of self rated health: cross sectional multi-country survey using anchoring vignettes. *BMJ* 2004; 328: 258.
- 73 Meijer RR. Diagnosing item score patterns on a test using item response theory-based person-fit statistics. *Psychol Methods* 2003; 8: 72–87.
- 74 King G, Wand J. Comparing incomparable survey responses: new tools for anchoring vignettes. *Polit Anal* 2007; 15: 46–66.